In order to find an explanation for individual reactions to transdermal glyceryl trinitrate (GTN) we studied the skin temperature and hemodynamic reactions in 63 healthy persons. The data were obtained before and after the application of GTN and Glycerin (GL) placebo patches, during one hour. The skin temperature was measured on both forearms, the local (left sided) and systemic (right sided) reaction on GTN was related to the skin fold and the calculated body fat content. The bilateral rise of skin temperature and its duration was higher and longer in obese than in lean persons mainly in obese women. The UV induced thermo and the later photothermo-reaction (Erythema) was reduced on the left forearm after the application of GTN and GL patches. The observed hemodynamic GTN effect confirmed known postural reactions, such as decreased arterial pressure ($\Delta$AP = -2.9%), increased heart rate ($\Delta$HR = +7.4%) and QTc prolongation ($\Delta$QTc = +4.9%) in upright position. An adverse drug effect with increased mean blood pressure ($\Delta$mAP = +12%) and increased heart rate ($\Delta$HR = +10.4%) mainly in supine position was observed in 11% of the participants, but only in men. Such a reaction was already described by Murell, 1879. Individual GTN effects were analyzed and related to habits and family history. In male smokers and in persons with hypertensive and diabetic close relatives, the hypotensive GTN effect was accentuated in supine position. In the upright position the group with hypertensives in the family presented a moderate hypotensive reaction without secondary tachycardia and the smokers presented only a slightly increased heart rate. Our observations suggest that individual reactions to transdermal glyceryl trinitrate (GTN) with its active component nitric oxide (NO) depends on physiological conditions, related to endogenous vasoactive substances, mainly the interaction with EDRF (the endogenous NO) and the activity of the Renin-Angiotensin System.


William Murell in 1879 observed that “nitroglycerine” given to friends, and other persons and also taken by himself in “six or seven minutes after an oral dose increased the pulse pressure, heart rate (HR) and the systolic arterial pressure (sAP) and the dicrotic notch became more prominent” (1). We believe that glyceryl trinitrate (GTN) named at that time as nitroglycerin, was tested in healthy men. This observation contrasts with the generally accepted concept that GTN in normal persons does not alter blood pressure and heart rate in supine position, but may provoke a postural hypotension with a secondary “reflex” tachycardia (2). With constant, low or high plasma nitrate levels, individual reactions are known: from abolished (3), attenuated (4) to opposite sympathetic hemodynamic drug effect with stimulation of the renin-angiotensin system (5). The introduction of GTN patches, using the transdermal route of application and thus avoiding the hepatic denitration of transmucosal absorption is an attempt to halt the development of nitrate tolerance which causes diminished effect on systemic blood pressure and HR (6). The development of nitrate tolerance and also the withdrawal syndrome depends on the continuous exposure to nitrates, but
METHODS

A total of 63 healthy volunteers (27 men and 36 women) were treated once with transdermal patches containing 10 mg of GTN. In a second turn, a total of 21 (9 men and 12 women) received a glycerin (GL) gauze patch as a placebo. Volunteers personal data are given as the mean ± standard deviation, for age: 34.8 ± 9.9 years; for height: 161.4 ± 9.8 cm; for body weight: 65.3 ± 11.9 kg; and for lean body weight (LBW): 51.9 ± 8.3 kg.

The LBW was calculated according to the formula: $E^2 \times R$, (E=height in cm; R for men = 0.002; R for women=0.0018) (10). The results allowed the calculation of the percentage of body fat content according to the formula:

\[
\frac{\text{Body weight (kg) - Lean body weight (kg)}}{\text{Body weight (kg)}}
\]

To understand the acute effect of transdermal GTN in healthy persons we analyzed the factors involved in drug absorption: the skin fold, body fat and skin temperature. Hemodynamic response to GTN was studied by the orthostatic reaction and related to individual variables as sex, subjective reactivity (as headache), smoking habit and family history of hypertension and diabetes in close relatives.

Skin fold was measured on both forearms insides over a distance of 3 cm; the mean value obtained was 9.6 ± 3.6 mm. Based on this parameter three subgroups were formed: Lean with 6.0 ± 1.8 mm, intermediate with 9.6 ± 3.6 mm and obese with 12.2 ± 4.4 mm.

In relation to smoking habits and hypertensive or diabetic close relatives, all the participants were classified as follows:

<table>
<thead>
<tr>
<th>Group</th>
<th>Smokers (more than 15 cigarettes/day)</th>
<th>Hypertension</th>
<th>Diabetes</th>
</tr>
</thead>
<tbody>
<tr>
<td>GTN treated (n=63)</td>
<td>24</td>
<td>39</td>
<td>21</td>
</tr>
<tr>
<td>GL treated (n=21)</td>
<td>12</td>
<td>11</td>
<td>8</td>
</tr>
</tbody>
</table>

Transdermal patches of GTN (Nitradisc®, Searle) of 10 mg and a gauze containing 5 ml of Glycerin (GL) were applied during 60 minutes on the left forearm insides, over an area of 15.9 cm² and 4.5 cm of diameter of GTN patch near the cubital region.

The following parameters were obtained: 1) The skin temperature was measured with an electrical thermometer in Celsius (°C) on both forearms before and immediately after drug application for one hour. Further measurements were taken just after a UV-B irradiation for five minutes (65 minutes) and finally after 125 minutes from the beginning of the drug application. The intensity of UV-B induced erythema on both arms was observed 24 hours later; 2) The hemodynamic parameters were studied in supine (10 minutes) and upright (2-3 minutes) position before and after the drug (GTN or GL) applications (60 and 125 minutes). The systolic (sAP) and diastolic (dAP) arterial blood pressures (mm-Hg) were registered by the auscultatory method and the mean arterial pressure (mAP) calculated by the equation:

\[
mAP = \frac{[(sAP - dAP) / 3] + dAP}{\text{HR} \times sAP \times 10^2}
\]

From the ECG (lead II), the HR (bpm) and the QT-interval (ms) were taken and corrected to QTc by Bazett formula (11).

The double product (DP) was calculated as DP = HR x sAP x 10².

Taken from a nomogram (Fig.1), one value was used to characterize the orthostatic ratio (OR) based on the difference between supine and upright positions for HR and mAP. A good hemodynamic adaptation on standing position reached values close to 100.
For values obtained “before” and “after” the transdermal application of GTN in combination of the variables of the subgroups, the statistical model used was one of the type “split-plot” (14). Significance of 5% is given for positive interaction “before” and “after” in all variables and their combinations.

For the skin temperature the repetition included eight levels due to significant interactions with intervals of confidence for multiple comparison by the Bonferroni Method (14).

RESULTS

GTN effect on skin temperature

Glycerin (GL) patches applied in 21 volunteers, on the left forearm did not alter the immediate (60 min.) and posterior (125 min.) skin temperature. The GTN patches in 63 participants, also applied during 60 min. on the left forearm caused a significant skin temperature elevation on both arms (Fig. 2).

The UV-induced immediate thermoreaction (skin temperature after 65 min.) and also the later photothermoreaction (Erythema after 24 hours) were attenuated after the previous application of GL and GTN patches on the left forearm.

Comparing the skin temperature on both forearms after 60 and 125 min. we observed that the left sided effect was distinct and prolonged.

The left sided skin temperature of persons with a 13.2 mm skin fold (n=17) was high both after 60 and 125 min. in 3.2 and 3.6% when compared with values obtained “before”. The right sided elevation of the skin temperature was smaller but also prolonged (2.6% after 125 min.). After the application of the GTN patches, obese persons with more than 17% of fat content, showed a distinct elevation of the left sided skin temperature. This observation was clearly seen in men; otherwise, obese women had a greater and longer elevation of the right sided skin temperature than obese men. (Table I)

GTN effect on hemodynamic-orthostatic reaction

A comparative study of GL and GTN effects revealed that Glycerin (GL) did not alter the arterial pressure, the heart rate (HR) and the QTc duration obtained from the ECG (Fig 3. and Table II).
Table I

Skin Temperature (°C)

<table>
<thead>
<tr>
<th></th>
<th>Left Forearm</th>
<th>Right Forearm</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>n</td>
<td>Before</td>
</tr>
<tr>
<td>Glycerin (GL)</td>
<td>21</td>
<td>32.3</td>
</tr>
<tr>
<td>Δ%</td>
<td></td>
<td>+0.4</td>
</tr>
<tr>
<td>Nitroglycerin (GTN)</td>
<td>63</td>
<td>31.1</td>
</tr>
<tr>
<td>Δ%</td>
<td></td>
<td>+2.3</td>
</tr>
<tr>
<td>Nitroglycerin Skin Fold Δ%</td>
<td></td>
<td>6.0±1.8 mm</td>
</tr>
<tr>
<td></td>
<td></td>
<td>9.6±3.6</td>
</tr>
<tr>
<td>Body Fat and Gender Δ%</td>
<td></td>
<td>13.2±4.4</td>
</tr>
<tr>
<td>&lt;12%♀</td>
<td>18</td>
<td>+0.9</td>
</tr>
<tr>
<td>12 - 17%♀</td>
<td>8</td>
<td>+2.1</td>
</tr>
<tr>
<td>&gt;17%♀</td>
<td>10</td>
<td>+2.5</td>
</tr>
<tr>
<td>&lt;12%♂</td>
<td>13</td>
<td>+1.4</td>
</tr>
<tr>
<td>12 - 17%♂</td>
<td>6</td>
<td>+2.4</td>
</tr>
<tr>
<td>&gt;17%♂</td>
<td>8</td>
<td>+3.4</td>
</tr>
</tbody>
</table>

(*) = Significance p<0.05

In all volunteers (n=63) a significant GTN effect appeared in the supine body position 60 minutes after the patch application with only a small decrease in the arterial pressure (-1.8%); in the upright position, GTN patches caused a significant decrease of the systolic (-3.1%) and diastolic (-3.2%) pressures and an increase in heart rate (+7.4%), when compared to “before”. Consequently the orthostatic reaction (OR) was reduced in 8%. The QTc duration from the ECG was prolonged from 0.4060±0.03 ms “before” to 0.4260±0.04 ms (p<0.05) “after” the GTN application, indicating a significant prolongation of 4.9% (Table III).

The prolongation of the QTc interval was also found in supine position, but only in men (n=27), regardless of complains of GTN induced headache.

Different groups presented significant altered orthostatic reaction (OR) in a range of -6.6 to -14.1% related to the situation “before”. The OR was negative (-8.0%) in the group with a positive family history of hypertension and diabetes and also in male smokers. In these groups the GTN patches provoked in the supine position a significant decrease in arterial pressure (sAP and dAP). In the upright position, only in the group with hypertensives in the family (n=39) and male smokers (n=15) we observed a clear hypotensive reaction. In this body position a tachycardial reaction but without hypotension was seen in the group with diabetes in close relatives.

Men (n=7) who presented headache after the the application of GTN patches during 60 min. responded differently: in the supine position, with a increased systolic pressure (+23%), with a decreased diastolic pressure (-10.4%) and a prolonged QTc interval (+5.4%); in upright position, only a decreased diastolic pressure (-10.4%) and a increased heart rate (HR) (+17.7%) was observed.
DISCUSSION

GTN effect on skin temperature

Only the GTN patches and not the GL (placebo) patches caused a significant elevation of the skin temperature on both arms. This observation suggests a local (left sided) and a systemic (right sided) vasodilating effect. The local as well as the initial (60 min.) and later (125 min.) GTN effect on skin temperature was accentuated when compared with the systemic effect.

Earlier studies had also shown that the sublingual nitroglycerin in normal subjects (n=11) increased the forearm blood flow, lowered the forearm vascular resistance due to a reduced venous tone. The mean arterial pressure (mAP) declined slightly (15).

An UV induced thermoreaction was used to differentiate alterations of the skin temperature due to capillary or vascular dilation. The UV induced erythema was evident on the right arm and pale or not visible on the left arm, indicating an inhibition of the local UV effect after previously applied GL and GTN patches. This observation may be related to a known inhibition of heparin activity due to glyceryl trinitrate (GTN) (16, 17). An interaction of GL and GTN with stimulants which releases heparin and also histamin may be considered. GTN as an inhibitor of heparin activity may also decrease the UV induced histaminergic reactivity in the skin.

Local factors may facilitate the drug absorption and influence the GTN effect. In obese men and women, the local and systemic drug effect on skin temperature is higher. In persons with a greater skinfold and a greater body fat content we observed a greater local, and a moderate systemic elevation of the skin temperature, when compared to lean persons. Approximately two hours (125 min.) after the application of the GL patches and UV application the skin temperature returned to the values obtained "before". After GTN patches and UV irradiation the skin temperature reached this level only in lean women and obese men; the GTN effect in these groups is abbreviated.

The transdermal absorption of GTN, a lipophilic compound, seems to be higher in obese persons. In obese women there may be a redistribution of the drug, as known to occur with barbiturates, which explains the prolonged effect on skin temperature not seen in lean women. A short drug effect was observed in obese men; a rapid breakdown and inactivation of the drug, or more likely, a counteracting vasoconstrictor GTN effect may be responsible for this effect.

Table II
Nitroglycerin – Hemodynamic parameters of groups

<table>
<thead>
<tr>
<th>Before / After</th>
<th>Supine Position (10 min)</th>
<th>Upright position (2 - 3 min)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Δ%</td>
<td>n</td>
<td>sAP</td>
</tr>
<tr>
<td>All persons</td>
<td>63</td>
<td>-1.8*</td>
</tr>
<tr>
<td>o F. hyp.</td>
<td>24</td>
<td>-4.4*</td>
</tr>
<tr>
<td>+ F. hyp.</td>
<td>39</td>
<td>-6.3*</td>
</tr>
<tr>
<td>o F. diab.</td>
<td>42</td>
<td>-8.3*</td>
</tr>
<tr>
<td>+ F. diab.</td>
<td>21</td>
<td>-2.1*</td>
</tr>
<tr>
<td>o Smokers</td>
<td>39</td>
<td>-4.8*</td>
</tr>
<tr>
<td>+ Smokers</td>
<td>24</td>
<td></td>
</tr>
<tr>
<td>o Headache</td>
<td>45</td>
<td>-2.1*</td>
</tr>
<tr>
<td>+ Headache</td>
<td>18</td>
<td>-12.0*</td>
</tr>
<tr>
<td>o Headache</td>
<td>20</td>
<td>-4.8*</td>
</tr>
<tr>
<td>+ Headache</td>
<td>7</td>
<td>+23.1*</td>
</tr>
</tbody>
</table>

sAP, dAP, mAP = systolic, diastolic, mean arterial blood pressure (mmHg); HR = heart rate (bpm); DP = double product; QTc = corrected systolic interval; OR = orthostatic reaction; Δ% = percent difference related to before values; (*) = significance p<0.05.
GTN effect on hemodynamic-orthostatic reaction

Our results obtained from 63 healthy middle aged persons confirm known acute GTN effects (18, 9). In the standing position, the decrease of the arterial blood pressure with increased heart rate (HR) resulted in a reduced performance of the orthostatic reaction (OR) not observed after application of GL patches. A negative orthostatic reaction (OR) indicates a moderate GTN effect with hypotension and tachycardia. The same result (-8%), as the mean value from all volunteers, was also observed in the group with a positive family history of hypertension, where in both body positions the arterial blood pressure was decreased but only in upright position the heart rate increased.

GTN increases the peripheral blood flow reducing the venous tone (15); with low GTN concentrations the venous dilatation predominates over that of arterioles (19).

A GTN induces postural hypotension due to “venous pooling”, enhances the activity of baroreceptors, leading to a secondary “reflex” change of the heart rate (HR) by a sympathetic discharge. Simultaneously an adrenergic stimulation of the left stellate ganglion, responsible for the “long QT syndrome” (20) may explain the QTc prolongation after GTN, as reported earlier (21). Supposing the stimulation of the left stellate ganglion due to GTN is correct and responsible for the QTc prolongation then the men studied, responded to GTN with an augmented sympathetic discharge.

In healthy persons, a previous individual physiological condition may explain the differences in the drug effect.

In the group with a positive history of diabetes in the family and also in male smokers, GTN patches provoked in the supine position, a clear hypotensive reaction and in upright position, a tachycardial response, without hypotension.

Individual and adverse responses to one small dose of transdermal applied GTN may be due to interactions of endogenous versus exogenous vasodilators, nitric oxide (NO) (22,23), endothelium derived relaxing factor (EDRF=NO) versus contracting factors (EDCF) (24) and endogenous vasoactive substances (25) and also the renin-angiotensin system (5,26).

Nitric oxide (NO) as an exogenous vasodilator, is derived from GTN, named nitroglycerin. GTN is metabolized in the presence of the sulfidryl group (SH) to nitric oxide (NO) in the vascular smooth muscle cells. The smooth muscle relaxing factor is cyclic guanosine 3’5’ monophosphate (cGMP) produced by the activation of the SH containing-enzyme, the guanylate-cyclase (27). The cGMP is the mediator for the exogenous NO and also the
The activation of cGMP by nitric oxide is competitive. The endothelium derived NO is an inhibitor of the GTN derived NO. A reduced EDRF release like under pathological conditions, fails to inhibit the exogenous NO activity on guanylate cyclase, thus potentiating the GTN effect. In healthy young people, due to a NO competition, the GTN derived NO activity may be reduced (25).

The interaction of the vasodilating GTN derived "exogenous" nitric oxide (NO) and the endothelium derived "endogenous" factor (EDRF=NO) is important but extrarenal renin release controlled by dietary potassium must be considered (28).

The renin-angiotensin system is involved in hemodynamic regulation as can be seen in the interaction of GTN and Captopril (29). Captopril, a compound containing a sulphhydryl group inhibits the angiotensin converting enzyme (ACE) and also the breakdown of bradykinin. Captopril reverts nitrate tolerance probably due to an activation of the SH-containing guanylate cyclase (30).

Low as well as a high renin activity mediated by angiotensin, modifies the adrenal cortex sensitivity leading to hyper- or hypotension (31). With a supposed reduced adrenal cortex sensitivity due to high renin activity, the GTN induced hypotension is accentuated as seen in male smokers and persons with hypertensive and diabetic close relatives.

We observed that the transdermal GTN provoked a mainly postural hypotension in 89% and a hypertensive reaction in 11% of all participants. In the same volunteers who participated in an earlier clinical study (32) and also in this study we observed low potassium (K⁺) level in plasma as well as an orthostatic hypodiastolic hypotension (33). The adverse paradoxal hypertensive GTN effect was only seen in men and was quite similar to the effect described by Murrel in 1879, after oral nitroglycerin doses (1). The men who responded to GTN with a headache had a tendency to an evident activity of vasoactive substances (serotonin, histamin, bradikinin and others) associated with a sympathetic reaction compatible with a "low renin hypertension". The greater group with a GTN induced postural hypotension may have an underlying "high renin" activity. Dietary potassium has also an influence over the renin release (28) and may be one of the determining factors for individual reactions to GTN (fig. 4). Counter regulatory neurohumoral factors and mechanisms such as enhanced catecholamin activity and stimulation of the renin angiotensin system may also be involved in the variability of nitrate tolerance (34).
**CONCLUSIONS**

The individual reactions to transdermal GTN revealed: 1) that one hour of application produces an effect on skin temperature mainly in obese women; 2) that the local and systemic GTN effect depends on arteriolar and not on (UV induced) capillary dilation; 3) that in healthy persons (n=63) transdermal GTN caused a moderate orthostatic reaction with a slightly decreased arterial pressure, increased heart rate and QTc prolongation; 4) that GTN in the supine position provoked individual reactions; in men (n=7) an adverse sympathetic reaction was observed associated with headache in supine and upright position.

**ACKNOWLEDGMENTS**

To M.T. Cavalcante, for technical assistance, to D.A. Botter and C. Qu. Aubin for the statistical analysis and to J.J. Leite for the revision of the paper.
RESUMO

Materiais e Métodos: Em 63 pessoas, antes e após aplicação tópica de discos com nitroglicerina® ou glicerina como placebo, foram avaliadas: a temperatura cutânea em ambos os antebraços e as relações hemodinâmicas em posição supina e ortostática.

Resultados: As respostas foram as seguintes: 1. Uma hora após aplicação unilateral (no antebraço esquerdo) houve aumento da temperatura cutânea também no antebraço direito, indicando efeito sistêmico; esse efeito era mais acentuado e prolongado em mulheres obesas. 2. A fototermo-reação (eritema) induzida com luz ultravioleta (UV) foi menor no antebraço esquerdo, sugerindo que a nitroglicerina age mais sobre aréolas do que sobre capilares do sistema vascular. 3. A aplicação transdérmica de nitroglicerina causou uma moderada reação ortostática com diminuição da pressão arterial e aumento da frequência cardíaca. A duração do QTc no ECG era prolongada. 4. Foram observadas as seguintes reações individuais em posição supina: em 7 homens houve uma reação adversa com aumento da pressão média em +12% e aumento da frequência cardíaca em +10%; a mesma reação (simpática?) foi descrito por Murell em 1879, após nitroglicerina v.o. Em homens fumantes e aqueles com hipertensão e diabetes em membros próximos da família, o efeito hipotensor após nitroglicerina transdérmica tinha sido acentuado em posição supina.

REFERENCES