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Differential effects of organic nitrates on arterial diameter among healthy Japanese participants with different mitochondrial aldehyde dehydrogenase 2 genotypes: randomised crossover trial

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ABSTRACT

Objectives: To determine whether polymorphisms at codon 487 (*1, GAA=Glu; *2, AAA=Lys) of mitochondrial aldehyde dehydrogenase 2 (ALDH2) influence nitroglycerine (glyceryl trinitrate (GTN))-induced vasodilation, and whether GTN or isosorbide dinitrate (ISDN) is a more effective antianginal agent in each ALDH2 genotype.

Design: A randomised, open-label, crossover trial with 117 healthy Japanese (20–39 years) whose genotypes were determined (*1/*1, n=47; *1/*2, n=48; *2/*2, n=22) was performed at Kyushu University Hospital, Fukuoka, Japan. Participants were randomly assigned to treatment: sublingual spray of GTN (0.3 mg) or ISDN (1.25 mg). After ≥1 week, measurements were repeated using the other drug. The main outcome measures were the maximal rate of increase in the brachial artery diameter determined by ultrasonography, the time required to attain maximal dilation (T_{max}) and the time required to attain 90% maximal dilation ($T_{0.9}$).

Results: The maximal artery diameter increase in response to GTN or ISDN did not differ among genotypes. However, GTN T_{max} was significantly longer for *2/*2 (299.7 s, 269.0–330.4) than *1/*1 (254.7 s, 238.6–273.4; p=0.0190). GTN $T_{0.9}$ was significantly longer in the *1/*2 (206.1 s, 191.7–219.3) and *2/*2 (231.4 s, 211.8–251.0) genotypes than *1/*1 (174.9 s, 161.5–188.3; p=0.0068, p<0.0001, respectively). In contrast, the time-course of ISDN-induced vasodilation did not differ among genotypes. GTN T_{max} and $T_{0.9}$ among *1 allele carriers (*1/*1 and *1/*2) were significantly shorter than those of ISDN, whereas the time course of GTN and ISDN vasodilation did not differ among participants carrying *2/*2.

Conclusions: The amplitude of GTN-induced vasodilation was not influenced by the ALDH2 genotype, but the response was significantly delayed in *2 allele carriers, especially *2/*2. GTN dilated the artery more quickly than ISDN in *1/*1 and *1/*2, but not in *2/*2.

ARTICLE SUMMARY

Article focus
- There were few reliable human studies on the influence of aldehyde dehydrogenase 2 (ALDH2) polymorphisms at codon 487 on the response to nitroglycerine (GTN). In particular, there was no information about the response to GTN in *2/*2 homozygotes.
- It was unclear whether GTN or isosorbide dinitrate (ISDN) is a more effective antianginal agent in each ALDH2 genotype.
- We aimed at comparing vasodilation induced by GTN and ISDN in individuals with different ALDH2 genotypes.

Key messages
- The maximal rate of increase in arterial diameter after GTN treatment did not differ among ALDH2 genotype groups.
- The time required to attain 90% maximal vasodilation was longer in the *1/*2 and *2/*2 groups than in the *1/*1 group.
- GTN (0.3 mg) induced vasodilation more rapidly than ISDN (1.25 mg) in the *1/*1 and *1/*2 groups, but there was no difference between GTN and ISDN in the *2/*2 group.

Trial registration number: UMIN000001492 (UMIN-CTR database).

INTRODUCTION

Nitroglycerine (glyceryl trinitrate (GTN)) has been the drug of choice to relieve angina pectoris since 1879 when William Murrell published his results on the clinical effect of GTN.[1] However, the effectiveness of GTN is not consistent among individual patients.[2] A sublingual 0.3 mg dose of GTN typically

ARTICLE SUMMARY

Strengths and limitations of this study

- The choice of healthy young individuals as participants may have strengthened the validity of the study; the participants' backgrounds were likely to be more homogeneous by excluding the influence of multiple concomitant factors that could not be controlled when studying patients or older subjects. However, the results could have been different if the study had been performed with patients with angina, because haemodynamic changes in patients might alter the response to nitrates.
- Sublingual spray formulations of nitrates were used instead of tablets to minimise the fluctuation in absorption rate. However, this may have made the dosage slightly unreliable.
- A randomised crossover design was used, and arterial diameter was determined by a single measurer blinded to genotype. However, the participants and investigators were not blinded to treatment.
- Arterial diameter was accurately measured using a semiautomatic ultrasonography system. However, the brachial artery diameter was measured instead of the coronary artery diameter, and moreover, we did not evaluate venous dilation.

alleviates chest pain, but some patients require a larger dose or repeated administration. The reason for this differential effect may be complex, since several biological processes are involved in mediating the pharmacological effects of GTN. GTN is a prodrug that must be reduced to glyceryl dinitrate to generate nitrite, which is further reduced to nitric oxide (NO). NO is the active metabolite that relaxes vascular smooth muscle by stimulating soluble guanylate cyclase to produce cyclic guanosine monophosphate.[3] [4] Proposed mechanisms for GTN bioactivation include the non-enzymatic interaction with thiol compounds such as cysteine and N-acetylcysteine,[5] and an enzymatic reaction such as glutathione S-transferase[6−8] and cytochrome P450[8] [9]; however, the relative importance of these mechanisms had not been understood until recently.

In 2002, Chen et al[10] identified mitochondrial aldehyde dehydrogenase 2 (ALDH2) as an important enzyme for GTN bioactivation.[11] They purified a protein fraction able to metabolise ^{14}C-GTN and sequenced its amino acids.[10] Subsequently, they showed that ALDH2 was essential for GTN metabolism using ALDH2 knockout mice.[12] They performed additional in vivo experiments in dogs to demonstrate the role of ALDH2 in GTN bioactivation.[13] These findings were supported by clinical studies. Mackenzie et al[14] reported that the effect of GTN administered by intra-arterial injection was 33% lower in healthy volunteers if the ALDH2 inhibitor disulfiram was given. The loss-of-function single nucleotide polymorphism at codon 487 of ALDH2, in which Glu (GAA, *1) is replaced with Lys (AAA, *2), is common in East Asian populations including Japanese. In an in vitro assay, the GTN reductase activity of ALDH2 *2 was shown to be two orders of magnitude lower than

that of the wild-type enzyme.[10] [15] Therefore, Mackenzie et al[14] evaluated the effect of GTN in 11 Asian *2 allele carriers (*1/*2 and *2/*2) and found that it was 40% lower than its effect in carriers of *1/*1. Li et al[16] reported that the effect of sublingual GTN was significantly lower in *2 allele carriers than in *1/*1 carriers among Chinese patients with angina; however, it is unclear whether this decrease in sensitivity to GTN should be considered in medical practice. Furthermore, the sensitivity to GTN of *2/*2 carriers should be distinguished from that of *1/*2 carriers, since the number of *2/*2 carriers included in previous studies was too small to be independently analysed. It is unknown whether another drug may be more effective for *2/*2 patients. In Japan, isosorbide dinitrate (ISDN) is the only alternative to GTN available for sublingual administration during anginal attacks. Since bioactivation of ISDN appears to be mediated by cytochrome P450 but not ALDH2,[9] [17] we wanted to compare the effectiveness of GTN and ISDN in individuals with different ALDH2 polymorphisms.

To answer these questions, we conducted a randomised crossover study to compare the vasodilation effect of GTN with that of ISDN in healthy young Japanese participants, including sufficient numbers of each ALDH2 genotype.

METHODS

Study design

From September 2009 to May 2010, we performed this randomised, open-label, crossover trial at Kyushu University Hospital, Fukuoka, Japan with volunteers whose ALDH2 genotypes were determined. After an initial treatment with a standard therapeutic dose of either GTN (0.3 mg) or ISDN (1.25 mg), the brachial artery diameter was continuously measured by ultrasonography. After ≥1 week, the participants received the other drug, and the measurements were repeated. The main outcome measures were the maximal rate of diameter increase, the time required to attain maximal dilation (T_{max}) and the time required to attain 90% of maximal dilation ($T_{0.9}$).

Participants

From April 2009 to March 2010, we recruited Japanese men and women (20−39 years old) among the students and staff of Kyushu University for participation in the study, and selected the final participants based on ALDH2 genotype. Persons who met the following criteria were excluded: (1) requiring treatment for a disease, (2) under medical treatment, (3) history of cardiovascular disease, (4) serious liver damage, (5) serious renal damage, (6) pregnancy, (7) history of adverse reaction to an organic nitrate, (8) receiving phosphodiesterase-5 inhibitors (sildenafil, vardenafil or tadalafil), (9) systolic blood pressure <90 mm Hg, (10) body weight <40 kg or >90 kg or (11) judged by the principal investigator (TS) to be inappropriate for participation in the study.

Genotyping

Genomic DNA was extracted from peripheral blood with a DNA isolation kit (GenTLE; Takara Bio, Ohtsu, Japan); for the 11 participants who did not wish to have blood drawn, DNA was extracted from saliva with a DNA self-collection kit (Oragene; DNA Genotek, Ottawa, Canada). DNA extraction was carried out with the kits according to the manufacturers' instructions. ALDH2 genotypes at codon 487 were determined by PCR and restriction fragment length polymorphism analysis. The PCR primer sequences were 5′-TTGGTGGCTACAA-GATGTCG-3′ (forward) and 5′-AAACACTGATGGCCT-CAAGC-3′ (reverse). The 324-bp PCR products were digested with *Tsp*RI (New England BioLabs, Ipswich, Massachusetts) at 65°C for 20 min. The restriction digest cut the *1 genotype DNA into three fragments (68, 255 and 1 bp) and the *2 genotype DNA into two fragments (323 and 1 bp), which were separated on a 3% agarose gel and visualised with ethidium bromide. Genotypes were determined by the existence of 255 bp and/or 323 bp fragments.

Interventions and outcome measures

Participants were prohibited from drinking alcohol and smoking cigarettes the night before testing, but meals were not restricted. After spending at least 60 min without exercise, participants were randomly assigned 1:1 to a single sublingual spray administration of GTN (0.3 mg; Myocor spray, Toa Eiyo, Tokyo, Japan) or ISDN (1.25 mg; Nitorol spray, Eisai Co, Tokyo, Japan) as initial treatment. A random allocation sequence was made by HA using random digits generated by the Mersenne Twister method[18] and conveyed to the investigators (SS, TY, HO and TS) by sealed numbered envelopes, one for each participant, with instructions to use the envelopes in numerical order. Participants rested at least 5 min in a supine position in a quiet examination room at 24°C, and then the right brachial artery was scanned in a longitudinal section 1 to 10 cm above the elbow in an arm maintained in the same position throughout the study, with a 10 MHz linear array transducer probe. To determine the maximal diameter of the artery, the end-diastolic diameter (length from one side of the media−adventitia interface to the other side) coincident with the R-wave on a monitoring electrocardiogram was continuously measured from 30 s before nitrate administration to 10 min after administration. This measurement was performed using a semiautomatic high-resolution B-mode ultrasound imaging system (UNEX EF 18G II, UNEX Corporation, Nagoya, Japan)[19 20] by a single expert (SS) blinded to genotype and recorded on a hard disk. Pulse rate and blood pressure were measured every 5 min. More than 7 days later, each participant received the other drug, and the measurements were repeated. The prespecified primary outcome measures were the maximal rate of increase in brachial artery diameter and the time required to attain maximal dilation (T_{max}). However, the time required to attain 90% maximal dilation ($T_{0.9}$) was also determined, so as to minimise fluctuation of the outcome measure. In addition, changes in pulse rate and blood pressure were assigned as secondary outcome measures.

Statistical analysis

The minimum sample size for this clinical trial was based on results obtained in previous studies. The mean and SD for the increase in forearm blood flow induced by GTN in *1/*1 and *2 allele carriers were reported by Mackenzie *et al.*[14] Because no data were available regarding the homozygous *2/*2 population, we estimated the mean and SD values based on an experiment by Chen *et al*[12] using ALDH2 knockout mice. According to their results, we assumed that the increase in blood flow in *1/*1 and *2/*2 genotypes was 4.5±3.36 and 1.9±1.99 (ml/100 ml/min), respectively. Power calculation indicated that the minimum sample sizes of *1/*1 versus *2/*2 carriers were 24:24, 27:19, 30:15, 37:13 or 43:11 to detect the difference (2.6 ml/100 ml/min) between these two genotypes with a statistical power of 90% and a p value of 0.05. The resultant sample size (*1/*1, n=47; *2/*2, n=22) had an 80% power to detect a minimal difference of 3.7% in arterial diameter change in response to nitrates between these groups. Likewise, there was an 80% power to detect a minimal difference of 35 s in time required for arterial dilation between the two genotypes.

Differences in baseline characteristics of participants were compared according to genotype, using the χ^2 test for the categorical variable and one-way analysis of variance (ANOVA) for continuous variables. Primary outcome measures were displayed according to genotype group using box-and-whisker plots. Overall group differences in the primary and secondary outcomes were determined by one-way ANOVA and analysis of covariance (ANCOVA) with sex, age, body mass index and systolic blood pressure as covariates. The Tukey−Kramer test was used for multiple comparisons. Differences in outcomes between GTN and ISDN were also determined by one-way ANOVA and ANCOVA with the same covariates in each genotype. All data were analysed with SAS version 9.2. A p value of <0.05 was considered significant.

RESULTS

The frequency of the ALDH2 *2 allele is approximately 20% in the Japanese population, although regional differences may exist.[21] The frequencies of each genotype are: *1/*1, 60% to 70%; *1/*2, 30% to 40%; and *2/*2, 3% to 5%. To overcome the difficulty of recruiting enough *2/*2 homozygotes by random screening only, we intentionally recruited participants who were unable to drink alcohol by advertisement at Kyushu University. We analysed genotypes of 305 individuals (*1/*1, n=156; *1/*2, n=111; *2/*2, n=38) and included 117 individuals in the trial to fulfil the required sample size (*1/*1, n=47; *1/*2, n=48; *2/*2, n=22). As shown in table 1, baseline characteristics did not differ among the genotype groups. We analysed the

Table 1 Baseline characteristics of the participants

	Total	*1/*1	*1/*2	*2/*2	p Value
No	117	47	48	22	
Male (%)	72.7	72.3	70.8	77.3	0.85
Age (years)	26.6±4.3	25.8±4.0	26.8±4.3	28.0±4.8	0.11
Height (cm)	167.5±8.4	168.0±8.5	168.1±8.1	165.0±8.9	0.41
Weight (kg)	59.6±9.3	59.9±8.5	59.9±10.4	57.7±8.4	0.62
Body mass index	21.2±2.3	21.3±2.1	21.1±2.7	21.1±1.9	0.91
Pulse rate (beats/min)	65.8±10.9	64.5±10.4	64.9±10.1	65.9±11.7	0.99
Systolic blood pressure (mm Hg)	119.5±11.8	118.4±13.5	117.0±12.5	118.3±10.9	0.60
Diastolic blood pressure (mm Hg)	67.1±8.4	66.7±8.7	65.9±8.1	64.3±6.8	0.42

Demographic and physiological parameters of the participants were evaluated by χ^2 test for sex and by one-way ANOVA for body mass index, pulse rate, systolic blood pressure and diastolic blood pressure. Results are expressed as mean±SD.

outcomes from all the participants without loss or exclusion.

Initially, we planned to assess the effectiveness of nitrates primarily with two parameters: (1) the maximal rate of increase in brachial artery diameter $(D_{max}-D_0)/D_0$, where D_0 is the diameter at time 0, and D_{max} is the maximal diameter achieved after nitrate administration; and (2) the time required to attain maximal dilation (T_{max}).

Unexpectedly, the first parameter $(D_{max}-D_0)/D_0$ did not differ among the genotype groups after either treatment (figure 1), even after adjusting for sex, age, body mass index and systolic blood pressure. This

finding suggests that GTN and ISDN eventually produce the same level of vasodilation regardless of genotype.

We then analysed the second parameter, T_{max}. As shown in figure 2, the GTN T_{max} was significantly longer in the *2/*2 genotype group than in the *1/*1 group (crude data, p=0.0190; adjusted data, p=0.0234), but no significant difference was observed between the *1/*1 and *1/*2 groups (crude, p=0.1390; adjusted, p=0.1834). The p trend among the three genotypes was significant for both crude (p=0.004) and adjusted (p=0.0002) data. In contrast, ISDN T_{max} did not differ among the genotype groups.
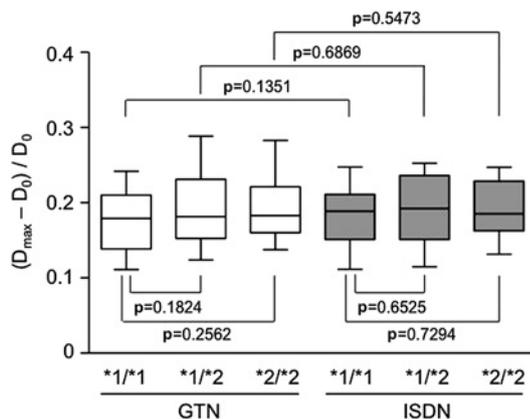

Figure 1 Maximal rate of increase in the brachial artery diameter. After sublingual administration of glyceryl trinitrate (GTN) (0.3 mg) or isosorbide dinitrate (ISDN) (1.25 mg), the brachial artery diameter was measured continuously for 10 min. The maximal rate of increase was calculated by the formula $(D_{max}-D_0)/D_0$, where D_0 was the baseline diameter, and D_{max} was the maximal diameter achieved. The values of $(D_{max}-D_0)/D_0$ after GTN treatment were: total, 0.188 (0.178−0.199); *1/*1 genotype, 0.176 (0.163−0.196); *1/*2 genotype, 0.195 (0.177−0.210); and *2/*2 genotype, 0.196 (0.172−0.221). The values of $(D_{max}-D_0)/D_0$ after ISDN treatment were: total, 0.187 (0.178−0.196); *1/*1 genotype, 0.183 (0.169−0.199); *1/*2 genotype, 0.189 (0.174−0.204); *2/*2 genotype, 0.189 (0.167−0.210). Results are shown as box-and-whisker plots indicating the 10th, 25th, 50th, 75th and 90th percentiles. Group results were compared by one-way ANOVA for crude data and by ANCOVA after adjusting for sex, age, body mass index and systolic blood pressure.
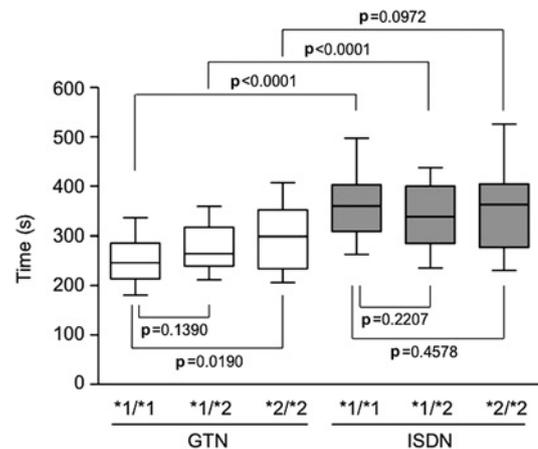
Figure 2 Time required to attain maximal dilation (T_{max}). After sublingual administration of glyceryl trinitrate (GTN) (0.3 mg) or isosorbide dinitrate (ISDN) (1.25 mg), the brachial artery diameter was measured continuously for 10 min. The values of the time when D_{max} was achieved (T_{max}) after GTN treatment were: total, 272.6 s (261.4−284.2 s); *1/*1 genotype, 254.7 s (238.6−273.4 s); *1/*2 genotype, 277.4 s (259.4−294.3 s); *2/*2 genotype, 299.7 s (269.0−330.4 s). T_{max} values after ISDN treatment were: total, 351.4 s (337.6−368.7 s); *1/*1 genotype, 365.0 s (340.5−389.9 s); *1/*2 genotype, 340.3 s (316.2−364.5 s); *2/*2 genotype, 347.0 s (320.7−391.3 s). Results for the different genotypes and nitrate treatments are shown as box-and-whisker plots indicating the 10th, 25th, 50th, 75th and 90th percentiles. Group results were compared by one-way ANOVA for crude data and by ANCOVA after adjusting for sex, age, body mass index and systolic blood pressure.

The brachial artery typically began to dilate about 1 min after nitrate administration. The diameter rapidly increased until 3 min, after which dilation slowed, and the diameter reached the maximum size at 4 to 6 min. However, it was often not easy to determine T_{max}, because the artery diameter usually approached the maximum size slowly. Thus, an artefact in the trace could significantly affect the determination of T_{max}. To avoid this problem, we introduced the parameter $T_{0.9}$, which is the time point at which the arterial diameter reached 90% of the maximal level, to minimise fluctuation of the outcome measure.

As shown in figure 3, GTN $T_{0.9}$ was significantly longer in participants with *2/*2 (crude, p<0.0001; adjusted, p=0.0002) and *1/*2 (crude, p=0.0068; adjusted, p=0.0157) genotypes than those with *1/*1. The p trend among the three genotypes was also significant (crude, p<0.0001; adjusted, p<0.0001). On the other hand, there was no difference in the $T_{0.9}$ of ISDN. A similar result was obtained when the time required for 50% maximal dilation was analysed (supplementary online figure).

Next we investigated which organic nitrate was more effective in each ALDH2 genotype. As shown in figure 1, the effects of GTN and ISDN on $(D_{max}-D_0)/D_0$ did not differ in any genotype group. In contrast, the T_{max} of GTN was significantly smaller than that of ISDN in the *1/*1 group (crude, p<0.0001; adjusted, p<0.0001) and *1/*2 group (crude, p<0.0001; adjusted, p<0.0001), but

no difference was observed between the effects of these nitrates in the *2/*2 group (crude, p=0.0972; adjusted, p=0.1023) (figure 2). The $T_{0.9}$ of GTN was also significantly smaller than that of ISDN in the *1/*1 group (crude, p<0.0001; adjusted, p<0.0001) and *1/*2 group (crude, p=0.0039, adjusted, p=0.0013), but this value did not differ between the treatments in the *2/*2 group (crude, p=0.1996, adjusted, p=0.3122) (figure 3).

The results of secondary outcome measures are summarised in the supplementary online table. No significant differences were observed in pulse rate, diastolic blood pressure or systolic blood pressure measured 10 min after nitrate administration among the ALDH2 genotype groups. Further, no serious adverse events were observed throughout the study.

DISCUSSION

Principal findings

First, the maximal rate of increase in diameter after GTN treatment did not differ among ALDH2 genotype groups, suggesting that GTN eventually induces maximal vasodilation regardless of genotype. Second, the GTN T_{max} was significantly longer in the *2/*2 group than in the *1/*1 group, and GTN $T_{0.9}$ was significantly longer in the *1/*2 and *2/*2 groups than in the *1/*1 group, suggesting that the *1 allele is associated with a more rapid response to GTN. However, this differential response was not observed after ISDN treatment. Third, GTN (0.3 mg) induced vasodilation more rapidly than ISDN (1.25 mg) in the *1/*1 and *1/*2 groups, but the T_{max} and $T_{0.9}$ values of GTN and ISDN did not differ in the *2/*2 group.

Strengths and limitations of the study

This study was carefully designed and conducted; however, our results require careful interpretation. First, the study participants were not patients with angina. Healthy, young volunteers were used because organic nitrates can also induce vasodilation in healthy people. The choice of participants may have strengthened the validity of the study; the participants' backgrounds were likely to be more homogeneous by excluding the influence of multiple concomitant factors that could not be controlled when studying patients or older subjects. In addition, using healthy, young participants made it easy to recruit a large enough sample of *2/*2 homozygotes. However, the results could have been different if the study had used patients with angina, because haemodynamic changes in patients might alter the response to organic nitrates. Second, we used sublingual spray formulations instead of tablets for both nitrates to minimise the fluctuation in absorption rate. However, this may have made the dosage slightly unreliable,[22] although this would have affected all the participants similarly and therefore should not have biased the results. Third, to minimise bias, a randomised crossover design was used, and arterial diameter was determined by a single measurer blinded to genotype. However, the participants and investigators were not blinded to
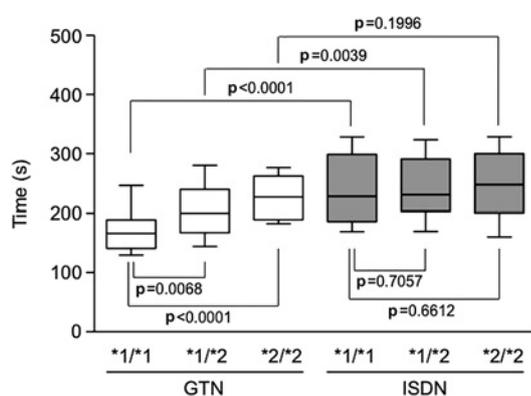
Figure 3 Time required to attain 90% maximal dilation ($T_{0.9}$). After sublingual administration of glyceryl trinitrate (GTN) (0.3 mg) or isosorbide dinitrate (ISDN) (1.25 mg), the brachial artery diameter was measured continuously for 10 min. The values of the time at which the diameter achieved 90% D_{max} ($T_{0.9}$) after GTN treatment were: total, 198.3 s (189.0−207.6 s); *1/*1 genotype, 174.9 s (161.5−188.3 s); *1/*2 genotype, 206.1 s (191.7−219.3 s); *2/*2 genotype, 231.4 s (211.8−251.0 s). $T_{0.9}$ values after ISDN treatment: total, 242.0 s (231.6−254.2 s); *1/*1 genotype, 236.8 s (220.8−257.0 s); *1/*2 genotype, 244.7 s (227.0−262.5 s); *2/*2 genotype, 247.4 s (221.5−273.3 s). Results for the different genotypes and nitrate treatments are shown as box-and-whisker plots indicating the 10th, 25th, 50th, 75th and 90th percentiles. Group results were compared by one-way ANOVA for crude data and by ANCOVA after adjusting for sex, age, body mass index and systolic blood pressure.

treatment. Fourth, arterial diameter was accurately measured using a semiautomatic ultrasonography system.[19 20] However, the brachial artery diameter was measured instead of the coronary artery diameter. Moreover, we did not evaluate venous dilation, although venodilation is thought to be a mechanism that is as important as coronary artery dilation for the antianginal effects of nitrates.[3 23 24]

Comparison with other studies

Other than the present study, three previous clinical studies have investigated the influence of the ALDH2 polymorphism on GTN effects, but the study performed by Mackenzie et al[14] was the only experimental study (clinical trial). In that study, GTN was injected into the brachial artery of healthy participants, and forearm blood flow measured by venous occlusion plethysmography was 33% lower if the ALDH2 inhibitor disulfiram was administered. Analysing 16 healthy East Asian participants (*1/*1, n=5; *1/*2, n=9; *2/*2, n=2), they also showed that the effect of GTN was 40% lower in carriers of the *2 allele than in participants with the *1/*1 genotype. However, the analysis did not differentiate between *1/*2 and *2/*2 genotypes, probably because the sample size was too small. In contrast, the present study included enough participants to allow analysis of the *2/*2 population. Moreover, we used ultrasonography to assess arterial dilation, which is a more direct method than plethysmography. The other two studies were both observational studies. Li et al[16] divided 80 Chinese patients with stable angina according to whether pain relief, assessed using a pain judgement scale, was obtained within 10 min after sublingual administration of GTN. ALDH2 genotypes of each group were then analysed, revealing that the effect of GTN was significantly smaller in carriers of the *2 allele than in participants with the *1/*1 genotype. Zhang et al[25] reported that in 142 Chinese patients with coronary heart disease, the rate of efficacious response to GTN was significantly higher in patients with the *1/*1 genotype than in those carrying the *2 allele. However, these studies did not distinguish between the *1/*2 and *2/*2 genotypes. Moreover, the efficacy of GTN was assessed according to the patients' subjective reporting of symptoms, potentially introducing bias. Therefore, few reliable human studies on the influence of the ALDH2 polymorphisms on the response to GTN were available until our study was conducted; in particular, there was no information about the response to GTN in *2/*2 homozygotes.

Based on the earlier studies, we initially hypothesised that the maximal response to GTN would be reduced in *2 allele carriers, but the results obtained were different from our expectation. Although the time required to obtain the response was different among the genotype groups, there was no difference in the size of response (the reason for this is discussed later). However, we cannot specify the reason for the difference in the results between earlier studies and ours. Probably multiple factors may be involved, such as study design, sample size, GTN dosage, studied vascular beds, measurement method or outcome measures.

Meaning of the study

The results of this study may have clinical implications for the daily use of organic nitrates. First, although there was no significant difference in the magnitude of response to GTN, the delay in attaining near maximal dilation was about 1 min for *2/*2 participants who took 0.3 mg of GTN. Given that the purpose of GTN is to provide rapid relief for chest pain, we believe that a 1 min delay cannot be ignored, although we observed only brachial artery dilation but not disappearance of the symptom. To practise personalised medicine for patients with angina, it may be useful to know the individual ALDH2 genotypes before treatment. If gene analysis is not available, a patient's genotype could be estimated by asking about drinking history or using the ethanol patch test.[26] Second, this study suggests the first choice drug for treating angina in terms of ALDH2 polymorphisms. Assuming that a rapid response is better, GTN may still be the first choice for *1/*1 patients and perhaps also for *1/*2 patients, because GTN dilated the artery more rapidly than ISDN in participants with these genotypes. However, ISDN may be a better choice for *2/*2 patients, because it dilated the artery as fast as GTN; moreover, larger or repeated doses of GTN often induce tolerance and oxidative stress, although ISDN can also cause tolerance.[27] In addition, this study may also be informative for basic research on organic nitrate pharmacology. The maximal rate of diameter increase was comparable among the genotypes. One explanation for this finding is that even the small amount of NO generated by ALDH2 *2 allele is sufficient for maximal response, because the GTN reductase activity of ALDH2 *2 is small but not nil.[15] Previously, maximal relaxation of rat aortic rings was shown to occur at only 3.4% of the maximal levels of cyclic guanosine monophosphate obtained with NO.[28] Another, perhaps more likely explanation is that enzymes other than ALDH2, such as glutathione S-transferase,[6 7 29] cytochrome P450[8 9] and xanthine oxidoreductase,[30] are also involved in the bioactivation of GTN. It has been suggested that ALDH2 activates GTN at the low concentrations achieved by therapeutic doses, whereas cytochrome P450 activates GTN at higher concentrations.[27] However, our results suggest that ALDH2 and other enzymes are concurrently involved in GTN bioactivation, even at low concentrations achieved by the standard therapeutic dose. The other enzymes involved may be able to generate a sufficient amount of NO without ALDH2, whereas ALDH2 specifically may be responsible for the rapid generation of NO.

Unanswered questions and future research

First, this study suggested that decreased ALDH2 activity delayed GTN-induced vasodilation in *2 allele carriers; however, direct evidence for delayed GTN bioactivation in *2 allele carriers has not been obtained. Therefore, pharmacokinetic studies evaluating serum

concentrations of GTN and its metabolites are needed to specifically assess whether GTN metabolism is delayed in *2 allele carriers. Second, we should also evaluate GTN-induced venous dilation in *2 allele carriers, since an in vitro experiment showed that venodilation was attenuated by pharmacological inhibition of ALDH2.[31] Third, the influence of the ALDH2 polymorphism on the action of GTN should be investigated with a clinical trial of patients with angina. No clinical trial has been conducted with patients so far; however, the study protocol should be carefully designed to enable precise determination of outcome measures. Fourth, basic studies are needed to identify the enzymes other than ALDH2 involved in GTN metabolism. Finally, it may be necessary to promote the development of a new medicine that is more beneficial than GTN.[32 33] This is not simply because the response to GTN varies among ALDH2 genotypes, but because repeated or long-term GTN administration may cause nitrate tolerance inactivating ALDH2 by superoxide generation,[4 27 34–36] and moreover, inactivated ALDH2 could disturb cardiovascular cell functions.[37 38]

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Competing interests None.

Ethics approval Ethics approval was provided by the Institutional Review Board for Clinical Trials, Kyushu University Hospital.

Contributors TS, KM, HA and FT-Y conceived and designed the study and wrote the protocol. SS, TY, HO and TS recruited participants, obtained informed consent and acquired the data. FS determined participants' genotypes. SS, TY, KM, and TS analysed and interpreted the data. HA undertook sample size calculation, randomisation, and statistical analyses. TS obtained funding. All authors wrote and revised the manuscript, and all approved the publication of the final version. All authors had full access to all the data in the study and take responsibility for the integrity of the data and the accuracy of the data analysis. TS is guarantor.

Provenance and peer review Not commissioned; externally peer reviewed.

Data sharing statement The full study protocol and data will not be publicly accessible. Interested individuals may contact the authors.

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CONSORT 2010 checklist of information to include when reporting a randomised trial*

Section/Topic	Item No	Checklist item	Reported on page No
Title and abstract			
	1a	Identification as a randomised trial in the title	1
	1b	Structured summary of trial design, methods, results, and conclusions (for specific guidance see CONSORT for abstracts)	2
Introduction			
Background and objectives	2a	Scientific background and explanation of rationale	5, 6
	2b	Specific objectives or hypotheses	6
Methods			
Trial design	3a	Description of trial design (such as parallel, factorial) including allocation ratio	7, 8
	3b	Important changes to methods after trial commencement (such as eligibility criteria), with reasons	No changes
Participants	4a	Eligibility criteria for participants	7
	4b	Settings and locations where the data were collected	7, 8
Interventions	5	The interventions for each group with sufficient details to allow replication, including how and when they were actually administered	8, 9
Outcomes	6a	Completely defined pre-specified primary and secondary outcome measures, including how and when they were assessed	9
	6b	Any changes to trial outcomes after the trial commenced, with reasons	9, 11, 12
Sample size	7a	How sample size was determined	9
	7b	When applicable, explanation of any interim analyses and stopping guidelines	Not applicable
Randomisation:			
Sequence generation	8a	Method used to generate the random allocation sequence	8
	8b	Type of randomisation; details of any restriction (such as blocking and block size)	8
Allocation concealment mechanism	9	Mechanism used to implement the random allocation sequence (such as sequentially numbered containers), describing any steps taken to conceal the sequence until interventions were assigned	8
Implementation	10	Who generated the random allocation sequence, who enrolled participants, and who assigned participants to interventions	8
Blinding	11a	If done, who was blinded after assignment to interventions (for example, participants, care providers, those	8

	11b	If relevant, description of the similarity of interventions	Not relevant
Statistical methods	12a	Statistical methods used to compare groups for primary and secondary outcomes	9, 10
	12b	Methods for additional analyses, such as subgroup analyses and adjusted analyses	9, 10

assessing outcomes) and how

Results

	13a	For each group, the numbers of participants who were randomly assigned, received intended treatment, and were analysed for the primary outcome	11
Participant flow (a diagram is strongly recommended)	13b	For each group, losses and exclusions after randomisation, together with reasons	11
Recruitment	14a	Dates defining the periods of recruitment and follow-up	7
	14b	Why the trial ended or was stopped	11
Baseline data	15	A table showing baseline demographic and clinical characteristics for each group	11, Table
Numbers analysed	16	For each group, number of participants (denominator) included in each analysis and whether the analysis was by original assigned groups	11
Outcomes and estimation	17a	For each primary and secondary outcome, results for each group, and the estimated effect size and its precision (such as 95% confidence interval)	11, 12, Figs. 1-3, Suppl. Table, Suppl. Figure
	17b	For binary outcomes, presentation of both absolute and relative effect sizes is recommended	
Ancillary analyses	18	Results of any other analyses performed, including subgroup analyses and adjusted analyses, distinguishing pre-specified from exploratory	11, 12
Harms	19	All important harms or unintended effects in each group (for specific guidance see CONSORT for harms)	12

Discussion

Limitations	20	Trial limitations, addressing sources of potential bias, imprecision, and, if relevant, multiplicity of analyses	13, 14
Generalisability	21	Generalisability (external validity, applicability) of the trial findings	14, 15
Interpretation	22	Interpretation consistent with results, balancing benefits and harms, and considering other relevant evidence	15, 16

Other information

Registration	23	Registration number and name of trial registry	2
Protocol	24	Where the full trial protocol can be accessed, if available	18
Funding	25	Sources of funding and other support (such as supply of drugs), role of funders	18

*We strongly recommend reading this statement in conjunction with the CONSORT 2010 Explanation and Elaboration for important clarifications on all the items. If relevant, we also recommend reading CONSORT extensions for cluster randomised trials, non-inferiority and equivalence trials, non-pharmacological treatments, herbal interventions, and pragmatic trials. Additional extensions are forthcoming: for those and for up to date references relevant to this checklist, see www.consort-statement.org.

Supplement Table. Changes in pulse rate and blood pressure induced by nitrates.

Pulse rate (PR), systolic blood pressure (SBP), and diastolic blood pressure (DBP) were measured at baseline and 10 min after glyceryl trinitrate (GTN) or isosorbide dinitrate (ISDN) administration. Results are expressed as percentage change and compared by one-way ANOVA.

		Total	*1/*1	*1/*2	*2/*2	P
GTN	%PR	0.21±9.32	0.13±10.73	-0.20±8.99	1.24±6.71	0.84
	%SBP	-5.69±5.49	-4.58±5.66	-6.46±5.41	-6.42±5.12	0.20
	%DBP	-13.83±9.16	-12.48±8.98	-14.43±8.62	-15.43±10.61	0.39
ISDN	%PR	1.33±7.45	1.73±8.83	0.10±6.71	3.12±5.48	0.26
	%SBP	-7.27±4.78	-7.13±4.88	-6.96±4.67	-8.23±4.93	0.57
	%DBP	-16.47±10.36	-17.86±10.45	-15.61±10.75	-15.43±9.42	0.51