Garlic Extract Diallyl Sulfide (DAS) Activates Nuclear Receptor CAR to Induce the Sult1e1 Gene in Mouse Liver

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Abstract

Constituent chemicals in garlic extract are known to induce phase I and phase II enzymes in rodent livers. Here we have utilized Car+/− and Car−/− mice to demonstrate that the nuclear xenobiotic receptor CAR regulated the induction of the estrogen sulfotransferase Sult1e1 gene by diallyl sulfide (DAS) treatment in mouse liver. DAS treatment caused CAR accumulation in the nucleus, resulting in a remarkable increase of SULT1E1 mRNA (3,200 fold) and protein in the livers of Car+/− females but not of Car−/− female mice. DAS also induced other CAR-regulated genes such as Cyp2b10, Cyp3a11 and Gadd45α. Compared with the rapid increase of these mRNAs levels, which began as early as 6 hours after DAS treatment, the levels of SULT1E1 mRNA began increasing after 24 hours. This slow response to DAS suggested that CAR required an additional factor to activate the Sult1e1 gene or that this activation was indirect. Despite the remarkable induction of SULT1E1, there was no decrease in the serum levels of endogenous E2 or increase of estrone sulfate while the clearance of oxoexogenously administered E2 was accelerated in DAS treated mice.

Results

Sult1e1 gene induction in mouse liver

Sult1e1 gene expression was analyzed in wild type and Car null female mice treated with DAS and DADS as described in the methods section. DAS dramatically induced (250-fold) the gene

Introduction

Garlic has been consumed for food and medicinal purposes worldwide for thousands of years. Garlic’s beneficial effects on human health have been known for a long time. Currently, the garlic plant itself as well as its numerous extracts are commercially available as dietary supplements. Epidemiological studies suggest garlic consumption has preventive effects for some types of gastric cancer [1–3]. The pharmacological basis for these preventive effects of garlic consumption have been suggested to be due to the inductions of various hepatic drug-metabolizing phase I and phase II enzymes [4–8]. Many of these enzymes are also induced by various CAR activators such as phenobarbital, phenytoin and TCPOBOP [9–13]. Whether CAR is essential for garlic extracts to induce these enzymes, however, has not been well established.

CAR, an orphan member of the nuclear steroid hormone receptor superfamily, was first found to trans-activate the CYP2B genes following phenobarbital and TCPOBOP treatment [14,15]. Subsequent investigations have characterized CAR as a xenobiotic-activated nuclear receptor, extended CAR activators to include numerous xenobiotics and endobiotics, and increased the large number of CAR-regulated genes [9,16–22]. As a result, the role of CAR has also widened from drug metabolism to various other hepatic functions such as energy homeostasis [23–26]. Upon xenobiotic activation, CAR is also known to become a risk factor that mediates the development of liver injuries such as steatosis, and hepatocellular carcinoma [27–30]. If, in fact, garlic activates CAR, this activation may have a wide range of both physiological and pathophysiological consequences.

Estrogen sulfotransferase (EST) encoded by Sult1e1 and SULT1E1 genes in mice and humans, respectively, is a phase II drug metabolizing enzyme localized in the cytoplasm [31,32]. Biochemical and enzyme kinetic analysis have shown that ESTs in mice and humans catalyze the conjugation of a sulfate group at the 3-hydroxyl position of estrogens at their physiological concentrations [33–35]. Since sulfated estrogens do not bind the estrogen receptor as effectively, EST is considered to be a major factor for estrogen inactivation and metabolism [31,32]. In fact, gene targeting disruption of Sult1e1 caused elevated estrogen levels systemically and in the amniotic fluid in pregnant mice [36]. On the other hand, sulfated estrogen circulating in the blood can be converted to estrogen at the target tissues by steroid sulfatase. Thus estrogen sulfotransferase works not only for elimination of estrogen but also to produce precursors for active estrogen in a form that is more soluble and easy to deliver to the peripheral tissues [31,32].

Here we have found that DAS, a major component of garlic extract, induced the estrogen sulfotransferase Sult1e1 gene in mouse livers and we have investigated the role of CAR in this induction using Car−/− mice. In addition, serum estrogen levels and estrogen clearance were measured to determine whether SULT1E1 induction by DAS modulated the plasma levels of estrogens.

Methods

Sult1e1 gene expression was analyzed in wild type and Car null female mice treated with DAS and DADS as described in the methods section. DAS dramatically induced (250-fold) the gene
expression at 24 hrs after DAS gastric administration (Fig. 1a). In contrast, only a marginal induction (7.3 fold) of the same gene was observed in Car null mice. Conversely, diallyl disulfide (DADS) induces this gene 1.9 fold in wild type mice and 15.5 fold in Car null mice, suggesting that this chemical does not activate CAR for Sult1e1 gene induction. In agreement with mRNA expression, SULT1E1 protein in liver cytosol was induced by DAS as shown in the western blot in Fig. 1b, left panel, while in Car null mice livers, DAS had negligible effects on the protein content. The protein detected by the anti-SULT1E1 antibody co-migrated with recombinant mouse SULT1E1 protein on the SDS-PAGE gel. The estrogen sulfotransferase activity of the liver extracts used in Fig. 1a is shown in Fig. 1b right panel. These results were well consistent with the detected mRNA levels and protein levels in Fig. 1a and Fig. 1b. Next, we analyzed changes in CAR protein content in the nucleus of liver cells following DAS treatment, since

**Figure 1. CAR dependent Sult1e1 gene induction.** (A) Wild type and Car null mice were given DAS (80 mg/100 g body weight) or DADS (8 mg/100 g body weight) by gastric gavage. 24 hrs after chemical administration, Sult1e1 and Car gene expression was analyzed by quantitative PCR as described in the methods section. The relative mRNA amount of each gene was calculated so that wild type vehicle treated samples have one unit of expression. (B) Left panel; Liver cytosol extracts from the same mice used in (A) were prepared and SULT1E1 protein content was analyzed by western blotting. Recombinant mouse SULT1E1 protein was used as positive control and indicated with R, while V represents vehicle treatment. Anti Hsp70 western blotting is shown as a loading control. Right panel; Estrogen sulfotransferase activities were measured for liver extracts from the same mice used in (A). n.d. stands for no activity detected. (C) Liver nuclear extracts from the same mice used in (A) were prepared and CAR protein content was analyzed by western blotting. Anti Laminβ1 western blotting is shown as a loading control.

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CAR is known to migrate into the nucleus from the cytoplasm upon its activation. Thus, we analyzed CAR protein levels in mouse liver nuclear extracts before and after DAS administration. We observed that levels of CAR protein in the nucleus were increased by DAS and DADS administration in wild type animals, while no CAR protein was detected with the same antibody in \textit{Car} null mice nuclear extracts. The effect of DAS was so dramatic that the CAR protein content in the nucleus was much higher than that of PB treated mice. The reason why DADS activates CAR for nuclear translocation but not for Sult1e1 gene activation may be an interesting topic for research in the future.

Gene inductions in DAS treated wild type and \textit{Car} null mice

In DAS treated mouse liver, other xenobiotic metabolizing enzymes were also induced as shown in Fig. 2. The most extensively analyzed CAR dependent gene in mouse liver, \textit{Cyp2b10}, was markedly induced almost to the same extent (216 fold) as \textit{Sult1e1} while no induction of this gene was observed in \textit{Car} null mice. In the case of \textit{Cyp3a11} and \textit{Gadd45b}, significant induction was observed in wild type mice while less induction was found in \textit{Car} null mice. For \textit{Cyp1a1}, no gene induction was observed in \textit{Car} null mice while significant induction was observed in wild type mice, suggesting that CAR plays a critical role in the induction of \textit{Cyp1a1} by DAS. These observations suggest that there may be regulatory elements in the mouse \textit{Cyp1a1} gene that can be activated by CAR, as in the case of the human \textit{CYP1A1} gene [37].

Time course \textit{Sult1e1} gene activation by DAS

Next, we analyzed the time course of the increase of gene expression after DAS treatment. \textit{Sult1e1} and \textit{Cyp2b10} gene expression was analyzed at 6, 24, 48, 72, and 96 hrs after DAS gavage (Fig. 3a). \textit{Cyp2b10} was very quickly induced compared to \textit{Sult1e1}. At 6 hrs, \textit{Cyp2b10} induction was already at the maximum while \textit{Sult1e1} induction was very low at this point. \textit{Sult1e1} expression levels increased up to 3000 fold and then decreased to an almost basal level at 72 hrs. In contrast, \textit{Cyp2b10} expression was gradually decreased from its 6 hr maximum (110 fold) to 60 fold at 48 hrs, and returned to basal levels at 72 hrs. In the meantime, we observed strong CAR accumulation in the nucleus at 6, 24, and 48 hrs while no CAR protein existed in the nucleus at 72 or 96 hrs (Fig. 3b). This dramatic difference in the time course for induction of these two genes suggests that partially independent mechanisms are playing roles for the induction of each gene. The time courses for induction of the \textit{Cyp3a11} and \textit{Gadd45b} genes were intermediate to those of \textit{Sult1e1} and \textit{Cyp2b10}, with maximum expression at 24 hrs and decreasing gradually to the basal level at 72 hr or 96 hrs.

![Figure 2. Gene activation by DAS.](https://example.com/figure2.png)

The expression of \textit{Cyp2b10}, \textit{Cyp3a11}, \textit{Gadd45b} and \textit{Cyp1a1} genes was analyzed in DAS treated and vehicle treated mice livers. The same RNA samples treated with vehicle or DAS from wild type and \textit{Car} null mice liver used in FIG. 1A were analyzed to determine the expression of these genes.

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Serum estrogen levels in DAS treated mice

We analyzed serum estrogen levels in DAS-treated mice. Surprisingly there was slight increase in estrogen levels in the serum of 4 week old mice after DAS treatment (Fig. 4a). Estrone sulfate levels in serum did not change with the same treatment (Fig. 4b). Thus we measured the clearance of exogenously administrated estrogen in ovariectomized mice with or without DAS pretreatment for 42 hrs to allow for maximal induction of the \textit{Sult1e1} gene. Then, the mice were administrated with estrogen (0.5 mg/kg) and estrogen levels in serum were determined at 3, 8, and 24 hrs after the administration (Fig. 4c). E2 was shown to be very rapidly cleared from the serum and at 24 hrs following administration, E2 levels in DAS treated and control mice were the same as the basal levels found in non E2 administered mice. At 3 hrs after E2 administration, the E2 levels in DAS treated mice were significantly lower than those of non treated mice, while at 8 hrs no difference was observed.

Discussion

Naturally-occurring organosulfur compounds including DAS found in garlic and onions can induce diverse phase I and phase II enzymes in mice and rats \cite{4,5,8}. Many of these enzyme inductions seem to be mediated by CAR activation \cite{6,7,38}. These gene inductions affect endogenous and exogenous chemical metabolism and the gene expression and metabolism change brought by garlic consumption should explain garlic’s beneficial effect on the body. Epidemiological studies established garlic’s preventive effects for colorectal and stomach cancer \cite{1–3}. Cancer incidents in other sites (head and neck, lung, breast, and prostate) are also suggested to be reduced with garlic consumption. Furthermore, \textit{Sult1e1} polymorphisms are shown to be a risk factor for breast and endometrial cancers \cite{39–41}. Thus, where \textit{CAR} and \textit{SULT1E1} genes are co-expressing, garlic may affect cancer formation and development by affecting \textit{SULT1E1} gene expression. However, there are no overall clear connections between the animal model studies and human epidemiological studies established so far. One group has shown that organosulfur compounds can reduce aflatoxin B1 genotoxicity in rats by inducing phase I and phase II enzymes that increase the metabolism of aflatoxin B1 \cite{4}. Since garlic extract can affect the expression of diverse genes thus resulting in many changes in biological processes, much additional work is necessary using human tissue derived model systems in order to establish the relevance of animal model findings on human health.

In this report we observed a very strong induction of the \textit{Sult1e1} gene in the mouse liver by gastrically injected DAS. CAR is a dominant factor in the induction of this gene, since only a marginal induction of this gene was observed in \textit{Car} null mice. Recently, mouse \textit{Sult1e1} was shown to be induced by many chemicals that activate \textit{CAR} \cite{11}. Among these chemicals, TCPOBOP, a specific mouse \textit{CAR} agonist, showed more potent...
induction in female mice than in males [11]. We are using female mice in this report and we observed slightly less Sult1e1 induction in male mice by DAS treatment (data not shown).

The Sult1e1 gene induction time course was found to be unique as shown in Fig. 3. Other canonical genes induced by CAR activation had their highest expressions between 6-24 hrs. In contrast, Sult1e1 gene induction was optimum at 48 hrs after chemical administration. This observation suggests secondary factors which are activated by CAR are involved in Sult1e1 gene induction. Recently, the paracrine regulatory mechanism of the SULT1E1 gene was characterized in HepG2 cells co-cultured with MMNK-1 cholangiocytes [42,43]. When HepG2 cells were co-cultured with CFTR siRNA treated MMNK-1 cells in a Transwell system, an unidentified factor secreted from MMNK-1 cells that is permeable through the Transwell membrane activated SULT1E1 gene expression in HepG2 cells. Whether such a paracrine mechanism is involved in Sult1e1 induction by DAS, thus causing a slow gene induction in contrast to other CAR responsive genes, remains unanswered at this point. Another recent report suggests that RORα suppresses Sult1e1 expression in mice [44]. The broad spectrum of RORα and RORγ in the regulation of phase I and phase II genes implies that these receptors have crosstalk with CAR. Whether such a crosstalk plays a role or not in Sult1e1 gene induction by DAS will be an interesting research target.

A striking finding of this report is that the strong Sult1e1 gene induction did not affect the endogenous E2 level while improved E2 elimination was observed when the hormone was exogenously administered to ovariectomized mice. A recent report demonstrated that E2 decreased in mice when the Sult1e1 gene was induced in mouse livers by glucocorticoid [45]. They suggested that one of the mechanisms by which glucocorticoid attenuates E2’s biological effect is augmenting Sult1e1 expression in the liver so that conversion of serum E2 to its sulfated form for eventual secretion from the body is accelerated. In contrast, although DAS treated mice have robust Sult1e1 induction (around 3200 fold), endogenous E2 was not affected although exogenous E2 clearance was faster as shown in Fig. 4. These results suggest that Sult1e1 in the liver may not be the dominant regulator for the E2 levels in the mice of our experimental systems. One additional point is that the endogenous E2 levels were analyzed in 4 week old female mice while 8 week old ovariectomized mice were used for E2 elimination experiment. Although the Sult1e1 gene in liver was induced in both cases (data not shown), we don’t know if this age difference can affect the whole body response to DAS. Moreover, DAS may have effect on estradiol synthesis pathways, estradiol turnover or expression/activity of SULT1E1 in the tissues other than liver. Further detailed pharmacokinetical, pharmacogenetical, and biochemical studies are necessary to determine why this strong induction of Sult1e1 does not alter serum E2 levels.

Materials and Methods

Animals

Car null mice with C3H/HeNCrI BR (C3H) genetic background were established as previously described [28]. Mice were kept in 12:12h dark/light cycle and fed with a standard solid diet ad libitum. C3H wild type and Car null female mice received DAS (80 mg/100 g body weight) and DADS (8 mg/100 g) via gavage using corn oil as the vehicle. Phenobarbital (1 mg/100 g) dissolved
in PBS was administrated intraperitoneally. Mice were sacrificed at specified hours after the administration of xenobiotics for liver RNA or serum preparation. All animal procedures were approved by the Animal Ethics Committee at NIEHS, NIH (Permit Number: ASP 00-19).

**Chemicals**

DAS (allyl sulfide, 97%) and DADS (allyl disulfide, 80%) were purchased from Sigma.

**Real Time RT PCR**

Total mouse liver (three C3H female mice for each data point) RNA was isolated with Trizol reagent (Invitrogen) and reverse transcription was performed using High Capacity cDNA Archive kits for the RNA. Quantitative real-time PCR was carried out with the 7900HT Fast Real Time PCR System (Applied Biosystems). For Nr1i3, FAM-CCATGAGCAGTATGCTAA-MGB probe was used with 5'-CCTGTGGCTAAGGAAACAG-3' and 5'-TCCTCAGTGGCCATGGTTTCTA-3' primers. For Sult1e1, Gypsy3a1, Gadd45b and Gyp1a1, pre-synthesized probes from ABI, Mm00499178_m1, Mm00456591_m1, Mm00731567, Mm00435123_m1, and Mm00487218_m1 were used, respectively. For normalization of gene expressions among individual mice, Gapdh gene expressions determined with TaqMan rodent glycer-aldehyde-3-phosphate dehydrogenase control reagent (Applied Biosystems) were utilized.

**Western blotting**

Mouse liver cytosol and nuclear extracts from C3H female mice were prepared as described previously [46]. Individual mouse liver extracts and pooled nuclear extracts from 3 mice were used for western blotting. The protein concentration of extracts was determined using the Bio-Rad Protein Assay reagent. Thirty and 100 μg protein samples for nuclear extracts and cytosol extracts respectively were separated on a SDS-PAGE gel and transferred to PVDF membranes. Anti SULT1E1 and recombinant mouse SULT1E1 were described in the previous report [47]. Anti CAR monoclonal antibody developed by Perseus Proteomics (Tokyo, Japan) was purchased from R&D Systems (Minneapolis, MN). Anti HSP70 and anti lamin β were from BD Biosciences (San Jose, CA) and Santa Cruz Biotechnology (Santa Cruz, CA), respectively.

**Estrogen sulfotransferase activity**

Estrogen sulfotransferase activities of liver extracts were determined as described previously [47]. Briefly, liver extracts were incubated with 2 μM [3H] labeled estradiol and 100 μM 5'-Phosphoadenosine-5'-phosphosulfate in 200 mM Tris-acetate (pH 8.0) buffer containing 10 mM magnesium acetate. The samples were incubated at 37°C for 30 min and extracted with dichloromethane. The [3H] label in the aqueous phase was counted to measure sulfated estradiol.

**Serum estrogen**

Ovariectomized or non-operated female C3H mice (7 weeks old) and female C3H mice (4 weeks old) were obtained from Jackson Laboratory (Bar Harbor, ME). For E2 clearance analysis, ovariectomized mice were treated with DAS or corn oil vehicle via gavage. Forty two hours after DAS treatment, beta E2 (0.5 mg/kg) was administrated subcutaneously. At 3, 8, and 24 hrs after E2 administration, mice were sacrificed for liver RNA preparation while blood samples were collected from the inferior vena cava. For endogenous E2 levels in 4 week old mice, mice were treated with DAS or vehicle. Serum and RNA were prepared at 48 hrs after the treatment. For estrone sulfate determination, 7 weeks old female C3H mice were utilized for DAS or vehicle treatment (48 hrs). Serum was isolated with BD Microtainer serum separator tubes (Becton, Dickinson and Co., Franklin Lakes, NJ). Serum E2 and estrone sulfate concentrations were determined with DSL-4800 Ultra-Sensitive Estradiol RIA kit and DSL-5400 Estrone Sulfate RIA kit (Beckman Coulter, Brea, CA).

**Statistical analysis**

Statistical analysis was performed by Student's t test to compare the estradiol elimination difference between vehicle and DAS treated C3H female mice.

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**Author Contributions**

Conceived and designed the experiments: TS MN. Performed the experiments: TS WDG KV TSW RM MN. Contributed reagents/materials/analysis tools: TS WDG KV TSW RM MN. Wrote the paper: TS MN. We also thank Dr. E. James Squires for helpful discussions.

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