Prolonged administration of secoisolariciresinol diglycoside increases lignan excretion and alters lignan tissue distribution in adult male and female rats

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Limited information is available on lignan metabolism and tissue distribution between sexes and the effects of prolonged lignan exposure on tissue concentrations. In the present study, excretion and tissue distribution of lignans were compared after 1 d and 7 d administration of flaxseed lignan secoisolariciresinol diglycoside (SDG) in male and female rats. Sprague–Dawley rats were daily gavaged per os with $^3$H-SDG (3.7 kBq/g body weight (bwt)) and unlabelled SDG (5.3 μg/g bwt). Urine, faeces, serum and tissues (liver, kidneys, bladder, spleen, lungs, brain, thymus, heart, muscle, adipose, mammary gland, ovaries, vagina, uterus, testis, seminal vesicles, coagulating glands and ventral prostate) were collected at 0, 12 and 24 h after a single lignan dose or after the last dose of 7 d exposure. The sample total lignan content was measured as radioactivity by liquid scintillation counting. In both sexes, majority of radioactivity was excreted in faeces (40–83 %) and urine (1–2–5–2 %). $^3$H-SDG administration increased radioactivity in all tissues at all time points, and the levels were further increased after prolonged SDG exposure. Liver contained majority of the tissue lignans (48–56 %) in both sexes after both exposure regimens. After prolonged SDG exposure, the serum lignan concentrations had reached a plateau which was approximately 4-fold of that of acute exposure, whereas in both sexes, concentrations in skin and kidneys still increased, indicating tissue accumulation. After prolonged exposure, females had higher lignan concentrations in heart and thymus at all time points, demonstrating sex-related differences in lignan tissue distribution and the possibility for sex-specific treatment responses. These findings facilitate identification of target tissues for local lignan actions in vivo.

Lignans: Secoisolariciresinol diglycoside: Tissue distribution

Secoisolariciresinol diglycoside (SDG) is a common lignan found in foods, and especially flaxseed and flaxseed-containing foods are rich in SDG$^{1,2}$. Ingested SDG is metabolised to enterolignans such as enterodiol and enterolactone in both rats and human subjects$^{3–5}$. Epidemiological evidence suggests that consumption of diet rich in lignans may decrease the risk of some chronic diseases such as cancer at multiple sites including breast$^{6–8}$, prostate$^{9}$, thyroid$^{10}$, glioma$^{11}$, and stomach$^{12}$. In experimental animal models, ingested purified SDG has been shown to inhibit mammary, skin and colon cancers$^{13}$. In postmenopausal women with newly diagnosed breast cancer, flaxseed ingestion before surgical removal of the tumour increased apoptotic index and decreased c-erbB2 score of the cancer tissue$^{14}$. Accordingly, in prostate cancer patients, flaxseed consumption decreased proliferation and increased apoptotic indices of the tumours$^{15,16}$. These pilot studies indicate the potential of flaxseed as a neoadjuvant therapy for inhibiting cancer progression in patients. In vivo studies indicate that flaxseed lignan SDG and its metabolites mediate at least part of the anticarcinogenic effects of flaxseed$^{17–21}$.

Very little is known about the effects of prolonged lignan administration on tissue concentrations. We have shown previously that SDG metabolites are accessible to various tissues in female rats, and that prolonged SDG exposure increased the concentration of lignan metabolites in the liver and adipose tissues$^{22}$. However, in the earlier study, the radio-labelled SDG was administered to the animals only once, either at day 1 or at day 10 after feeding the animals unlabelled SDG. It is therefore unclear if the SDG metabolites accumulated in the tissues during the feeding period. There is also a lack of studies comparing the serum and tissue distribution of lignans in males and females. One human pharmacokinetic study performed with a single dose of SDG showed a sex difference in the time to reach the maximum lignan concentration in plasma and the mean residence time$^{23}$. We hypothesise that the SDG metabolites increase in the serum and accumulate in the tissues after prolonged exposure, and that their levels differ in males and females. Therefore, we determined the excretion and serum and tissue distributions of lignan metabolites after a single-dose and prolonged 7 d (one dose per day) regimen of flaxseed SDG in adult male and female rats. Identifying possible sex differences in lignan tissue distributions is important when assessing efficacy and safety of the compounds. A better understanding of SDG metabolism and lignan tissue distribution after acute

Abbreviations: i.p., intraperitoneal; s.c., subcutaneous; SDG, secoisolariciresinol diglycoside.

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and prolonged exposure is needed to identify the target organs for the lignan actions in vivo, and to determine the potential effects for different consumer groups of lignan-containing foods.

Materials and methods

$^{3}H$-labelled secoisolariciresinol diglycoside

SDG isolated from flaxseed$^{(22)}$ was labelled with $^{3}H$ into benzyl methylene groups of the molecule by Amersham International (Little Chalfont, Buckinghamshire, UK). Previous studies have shown that the label in this position is stable and not affected by intestinal metabolism$^{(22,23)}$. The radiochemical purity of $^{3}H$-SDG was 98.5%. The specific radioactivity of the product was 999 GBq/mmol as described previously$^{(22)}$.

Animals, diets and experimental design

Eight-week-old male ($n=18$) and female ($n=18$) Sprague–Dawley rats were obtained from Charles River (St Constant, PQ, Canada). An American Institute of Nutrition-93G-based diet$^{(24)}$ modified to contain 20% fat was used as a basal diet in the study. The high fat content was used to mimic a Western diet. The diet consisted of casein (200 g/kg), t-cystine (3 g/kg), sucrose (100 g/kg), maize starch (301.06 g/kg), dextrose (99.5 g/kg), soyabean oil (200 g/kg), $\alpha$-butylhydroquinone (0.04 g/kg), cellulose (50 g/kg), American Institute of Nutrition-93G mineral mix (35 g/kg), American Institute of Nutrition-93G vitamin mix (10 g/kg) and choline chloride (1.4 mg/kg). After 1-week acclimatisation to the high-fat American Institute of Nutrition-93G diet, the rats were weighed daily, and were gavaged accordingly per os once per day with $^{3}H$-SDG (3.7 kBq/g body weight) and unlabelled SDG (5.3 µg/g body weight) in 1 ml of distilled water. Serum and tissue samples (liver, kidneys, bladder, spleen, lungs, brain, thymus, heart, muscle, interscapular subcutaneous (s.c.) and ventral intraperitoneal (i.p.) adipose) were collected from all the rats after 1, 12 and 24 h after a single lignan dose and after the last dose of 7 d lignan administration (three rats per sex at each time point). The body weights of female and male rats were 214 (SE 4) and 352 (SE 4) g after 1 d exposure ($n=9$ for both sexes) and were 244 (SE 4) and 419 (SE 4) g after 7 d exposure ($n=9$ for both sexes). In rats, interscapular s.c. adipose tissue consists mainly of brown adipose, while i.p. adipose tissue is white. Additionally, testis, seminal vesicles (without secretion), coagulating glands and ventral prostate were collected from male rats, and mammary gland, ovaries, vagina and uterus were collected from female rats. The individual rat urine and faecal samples were collected from the metabolic cages. During the collection period, rats had access to water and basal diet ad libitum. The weight or volume of the collected samples was recorded, and the samples were stored at −20°C until analysis.

Sample preparation and liquid scintillation counting

The radioactivity of the samples, i.e. all lignan metabolites of administered $^{3}H$-SDG, was measured using an earlier described method$^{(22,23)}$ with slight modifications. Depending on the size and the fat content of the analysed tissue, the samples were dissolved in 0.5 or 0.7 ml of 1 M-hyamine hydroxide in methanol (9:1, v/v; Packard Bioscience B.V., Groningen, The Netherlands), and were incubated overnight in a shaking water bath at 45°C. A sample of dissolved tissues (50–100 µl), sample of faeces homogenised in distilled water (100 µl), or serum and urine samples (50 µl) were mixed with 5 ml of scintillant (Cytoscient ES, ICN Biomedicals, Costa Mesa, CA, USA). Duplicate samples were counted for 5-min periods using the TRI-CARB 2900 TR liquid scintillation analyser (Packard Instrument Company, Meriden, CT, USA).

The counting efficiencies in different tissues varied from 48 (spleen) to 77% (ovaries and uterus). The results were corrected for counting efficiency, possible losses in sample preparation, and chemical and colour quenching as described previously$^{(22)}$. The measured radioactivity in samples was converted to picomole equivalents of $^{3}H$-SDG by dividing it by the specific activity of the radioisotope. The percentage of radioactivity excreted in faeces and urine or recovered in tissues was calculated by dividing the 24-h total radioactivity excretion or recovery in tissues by the amount of gavaged radioactivity (the dose administered 24 h before sample collection) × 100. Specific tissue recovery of radioactivity was also estimated as a percentage of the sum of all tissue recoveries, i.e. specific tissue picomole value divided by the sum of picomole values of all measured tissues × 100.

For some of the organs with the highest recovered radioactivities, contributions of residual blood as percentage of tissue radioactivity were calculated according to the data published by Smith$^{(25)}$. The residual blood content values (ml/g of tissue) used for residual blood content in liver, brain, kidneys, heart and spleen were 0.182, 0.037, 0.209, 0.243 and 0.157, respectively$^{(25)}$. The proportion of serum was calculated to account for 65% of the rat blood volume$^{(26)}$.

Ethical approval of the study

Animal care and all experimental procedures in the study were approved by the University of Toronto Animal Ethics Committee, and were performed according to the Guide to the Care and Use of Experimental Animals$^{(27)}$.

Statistical analyses

The statistical analyses were performed using Statistica software for Windows (StatSoft, Tulsa, OK, USA). Because the distribution of the radioactivity data was normal as determined using the Shapiro–Wilk test, the differences in radioactivity content in different tissues were analysed with one-way ANOVA followed by post hoc Tukey’s test. The acceptable level of significance was set at $P\leq0.05$ for all the analyses.

Results

Excretion of lignans into urine and faeces

In both male and female rats, most of the radioactivity was excreted in faeces (40–83% of the administered dose) and urine (1.2–5.2% of the administered dose) during the 24-h period after both 1 d (a single dose) and 7 d (one dose...
Serum radioactivity

In both female and male rats, the serum $^3$H concentrations were significantly increased 12 h after a single $^3$H-SDG dose, and remained constant for 24 h (Fig. 2). The prolonged lignan exposure increased the serum radioactivity significantly in both male and female rats, and the levels remained constant during the 24 h period (Fig. 2). No differences in serum radioactivity levels were measured between sexes (Fig. 2).

Tissue distribution of lignans

After a single dose of $^3$H-SDG (1 d exposure), the radioactivity concentrations were similar after 12 and 24 h in all the measured tissues in both sexes except in the male bladder that contained higher $^3$H activity at 12 h time point (Fig. 3(A)). The 7 d prolonged SDG administration significantly increased the ligan concentrations in all tissues ($P<0.05$) compared with the 1 d exposure. In males and females, the highest $^3$H tissue concentrations were measured 12 h after the last $^3$H-SDG dose in muscle, thymus, spleen, liver, bladder, and brain. In addition, heart in males and s.c. adipose tissue and lungs in females had the highest radioactivity at 12 h. In males, the tissue $^3$H concentrations were high after 12 and 24 h in skin, kidneys and lungs, and in females, they were high in skin and kidneys. However, the radioactivity concentration remained constant over 24 h in i.p. adipose tissue. The highest radioactivity concentrations were measured in female heart tissue after 7 d $^3$H-SDG exposure ($P<0.05$), and they remained similar over the 24 h period (Fig. 3(B)). The lowest radioactivity concentrations were found in adipose tissues in both sexes ($P<0.05$) (Fig. 3). In all tissues, the radioactivity concentrations were lower than those in the serum.

Differences in tissue ligan distribution between males and females

Twelve hours after a single SDG dose, male bladder contained significantly higher ligan concentrations than female bladder (Fig. 3(A)). All other sex-related differences in tissue ligan distribution were observed after 7 d SDG exposure, and the concentrations were higher in females than in males. Females had higher tissue ligan concentrations in thymus and heart at all time points, while in lung tissue, the ligan concentrations were higher at 0 and 12 h (Fig. 3(B)). In female muscle tissue, the ligan concentration was higher than that in males at 0 h, and that in s.c. adipose tissues at 12 h (Fig. 3(B)). In all other measured tissues, the ligan concentrations were similar in both sexes.

Ligan distribution in female and male reproductive tissues

Similar to non-reproductive tissues, prolonged $^3$H-SDG administration significantly increased ($P<0.05$) the radioactivity concentrations in all female (Fig. 4) and male (Fig. 5) reproductive tissues. In females, the mammary gland had the lowest and the vagina had the highest tissue ligan...
concentrations \((P<0.05)\) (Fig. 4(B)). After prolonged exposure, the tissue \(^3\text{H}\) concentrations in ovaries, uterus and mammary gland remained similar over the 24 h period (Fig. 4(B)). In males, the highest radioactivity concentrations were observed in seminal vesicles 12 h after the last \(^3\text{H}-\text{SDG}\) dose of 7 d exposure (Fig. 5(B)). Testis, seminal vesicle and ventral prostate also had the highest radioactivity concentrations at 12 h, and coagulating glands had the highest radioactivity concentrations at 24 h after the last lignan dose (Fig. 5(B)).

Organ distribution of lignans

As expected, in both sexes, liver contained majority of the recovered tissue radioactivity after both a single-dose (53–56 % in males and 51 % in females) and multiple-dose (one dose per day for 7 d) administration (48–56 % in males and 50–55 % in females) (Table 1). However, significant proportions of the \(^3\text{H}\) activity were also recovered in brain and kidneys (8–16 and 6–11 %, respectively). In male reproductive organs, testes contained majority (6–11 %) of the recovered tissue radioactivity, and in females, uterus contained up to 1–2 % of the recovered tissue radioactivity (Table 1).

The percentage of radioactivity recovered of the administered \(^3\text{H}-\text{SDG}\) dose was relatively low in all organs (Table 1). Liver contained up to 2–1 % of the administered dose, while in all other tissues, the recovery was below 1 %. There were no significant differences in organ weights adjusted for the body weight of 1- and 7 d SDG-administered.
male or female rats (data not shown). At necropsy, no visible changes in tissue colour or texture were observed in rats exposed to SDG compared with non-exposed animals.

In individual samples of liver, spleen, kidneys, heart and brain, residual blood accounted for 13–22, 11–23, 15–28, 14–30 and 2–5 % of the measured radioactivity, respectively. No statistically significant differences in the tissue residual blood radioactivities between sexes or duration of lignan exposure were observed.

Discussion

SDG is known to be metabolised by human intestinal microbiota to secoisolariciresinol and then to enterolignans enterodiol and enterolactone (28,29) that are found in urine, faeces and serum of SDG-exposed rats and human subjects (3–5,30). The present study has shown for the first time increased tissue lignan concentrations in rats after prolonged (7-d) SDG exposure compared with acute (1-d) exposure regimen and lignan accumulation in specific tissues. Moreover, tissue-specific sex differences in lignan distribution were observed.

The majority of the radioactivity was excreted in urine and faeces after 1 d 3H-SDG administration, which was further increased after prolonged lignan exposure, in agreement with previous studies (22,31). In the present study, the urinary and faecal excretion of lignans was similar in males and females. However, while the faecal excretion was similar to, the urinary excretion of SDG-derived lignans at 24 h was about two times less than that in our previous study in female rats (22), but was similar to that in male rats (31). As in human subjects, the urinary excretion of lignans varies significantly between rats. The lignan excretion may vary few folds even between the littermates housed and treated in similar conditions and having the same basal diet (unpublished results from our laboratory). The 40 % urinary excretion observed in human subjects fed SDG extract (35) was higher partly because it was based on cumulative urinary excretion at up to 48–72 h instead of 24 h. The difference in urinary excretion of SDG in rats and human subjects or pigs may be explained in part by individual differences in lignan uptake, liver metabolism and biotransformation capability of ingested lignans by gut microbiota.

In the previous study (22), majority of faecal radioactivity was excreted during the first 12 h after a single 3H-SDG dose, while in the present study, a majority was excreted after 12 h. This may be due to the fact that rats in the previous study were fasted before SDG administration, which may have resulted in faster faecal excretion, while rats in the present study were not fasted. In 7 d exposed rats, majority of lignans were excreted in urine 12–24 h after the last SDG dose, indicating delayed excretion compared with acute exposure when most of the lignans were excreted during the first 12 h.

Prolonged exposure to SDG increased serum lignan levels approximately 4-fold in both male and female rats. This concurs with previous studies with flaxseed showing increased plasma and serum enterolignan concentrations in human subjects after prolonged flaxseed consumption (32,33). In rats in the present study, the serum lignan concentrations levelled off after 7 d SDG administration. Accordingly, in women consuming daily 25 g of flaxseed, no significant differences in the plasma enterolignan levels were observed within the eighth day of flaxseed consumption (32), indicating that approximately 1-week exposure to flaxseed or SDG is sufficient to stabilise the serum enterolignan concentrations. Moreover, the radioactivity concentrations in serum were
always higher than those in tissues. However, after 7 d 3H-SDG exposure, the tissue radioactivity concentrations remained elevated at 24 h time point compared with 0 h time point in skin and kidney in both sexes, and in s.c. adipose tissue, muscle, spleen, lung, brain, heart, seminal vesicles, testis and ventral prostate in males. The further increase in the tissue lignan concentrations after serum concentration has plateaued indicates slow tissue-specific lignan accumulation and the possibility for tissue concentrations higher than those in serum after long-term exposure. When sesaminol triglucoside, which like SDG can be metabolised to entero-
lignans, was administered to rats at a high dose (1500 mg/kg per d), caecal and colon tissue enterolignan concentrations exceeded those of plasma(34). Also in men, higher concentrations of enterolignans in prostate tissue than in plasma have been reported(35). These findings by others and the accumulation of SDG-derived lignans in specific tissues in the present study suggest an active transport mechanism of lignans. Thus, serum concentrations alone do not fully reflect the concentrations in specific tissues.

The specific metabolites of 3H-SDG were not measured in the present study, but our previous study(23) has shown that the radioactivity in urine after feeding 3H-secoisolariciresinol diglycoside (SDG). The data are expressed as means with their standard errors, n 3 rats per time point. * Mean values between panels with unlike superscript letters were significantly different between time points for a tissue (P≤0.05).

![Fig. 5. Male rat tissue radioactivity concentrations at different time points over 24 h after (A) a 1 d (single dose) and (B) 7 d (one dose per day) per os administration of 3H-secoisolariciresinol diglycoside (SDG). The data are expressed as means with their standard errors, n 3 rats per time point. * Mean values between panels with unlike superscript letters were significantly different between time points for a tissue (P≤0.05).](https://doi.org/10.1017/S0007114510001194)

In rats, s.c. adipose at interscapular region contains mainly brown adipose tissue, while i.p. adipose tissue is white. We found a difference in the lignan concentrations between s.c. and i.p. adipose tissues, indicating that lignan uptake or accumulation between different adipose types may vary. Our study indicated for the first time sex differences in total lignan concentration in specific tissues. After prolonged 3H-SDG exposure, higher radioactivity concentrations in heart, thymus, lung, muscle and s.c. adipose tissues were observed in female rats. Indeed, the radioactivity concentrations in heart and thymus were higher in females than in males at all time points measured. A sex difference in total
Table 1. Recovered tissue radioactivities at different time points
(Mean values with their standard errors)

<table>
<thead>
<tr>
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<th>Recovered tissue radioactivity %* (% radioactivity recovered of administered dose)</th>
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<tbody>
<tr>
<td></td>
<td>Day 1</td>
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<tr>
<td></td>
<td>12 h Mean (SE)</td>
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<tr>
<td></td>
<td>Mean SE</td>
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<tr>
<td>Male organs</td>
<td></td>
</tr>
<tr>
<td>Liver</td>
<td>56.4 (0.67) 3.6 (0.02) 53.3 (0.53) 1.5 (0.02)</td>
</tr>
<tr>
<td>Brain</td>
<td>8.5 (0.10) 0.5 (0.00) 8.9 (0.09) 0.6 (0.00)</td>
</tr>
<tr>
<td>Kidneys</td>
<td>9.0 (0.11) 0.8 (0.01) 8.8 (0.09) 0.3 (0.00)</td>
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<tr>
<td>Lungs</td>
<td>5.2 (0.06) 0.5 (0.01) 5.7 (0.06) 0.5 (0.00)</td>
</tr>
<tr>
<td>Heart</td>
<td>4.1 (0.05) 0.2 (0.01) 4.3 (0.04) 0.3 (0.00)</td>
</tr>
<tr>
<td>Spleen</td>
<td>3.7 (0.04) 0.3 (0.01) 3.5 (0.03) 0.6 (0.01)</td>
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<tr>
<td>Thymus</td>
<td>1.9 (0.02) 0.4 (0.00) 2.1 (0.02) 0.1 (0.00)</td>
</tr>
<tr>
<td>Bladder</td>
<td>1.4 (0.02) 0.3 (0.00) 0.5 (0.01) 0.1 (0.00)</td>
</tr>
<tr>
<td>Testes</td>
<td>6.1 (0.11) 3.1 (0.01) 9.3 (0.09) 0.2 (0.00)</td>
</tr>
<tr>
<td>Seminal vesicles</td>
<td>2.0 (0.04) 1.0 (0.01) 1.9 (0.03) 0.9 (0.00)</td>
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<tr>
<td>Ventral prostate</td>
<td>1.3 (0.02) 0.0 (0.00) 1.2 (0.01) 0.2 (0.00)</td>
</tr>
<tr>
<td>Coagulating glands</td>
<td>0.3 (0.01) 0.2 (0.00) 0.6 (0.01) 0.1 (0.00)</td>
</tr>
<tr>
<td>Female organs</td>
<td></td>
</tr>
<tr>
<td>Liver</td>
<td>51.2 (0.52) 1.3 (0.02) 51.3 (0.51) 1.9 (0.02)</td>
</tr>
<tr>
<td>Brain</td>
<td>1.3 (0.13) 0.5 (0.00) 14.1 (0.14) 1.3 (0.01)</td>
</tr>
<tr>
<td>Kidneys</td>
<td>10.5 (0.11) 0.4 (0.01) 9.5 (0.10) 0.3 (0.00)</td>
</tr>
<tr>
<td>Lungs</td>
<td>8.7 (0.90) 0.8 (0.01) 9.7 (0.10) 0.4 (0.00)</td>
</tr>
<tr>
<td>Heart</td>
<td>6.8 (0.07) 0.6 (0.01) 7.0 (0.07) 0.2 (0.00)</td>
</tr>
<tr>
<td>Spleen</td>
<td>4.1 (0.04) 0.8 (0.01) 3.3 (0.03) 0.2 (0.00)</td>
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<tr>
<td>Thymus</td>
<td>3.1 (0.03) 0.7 (0.01) 2.6 (0.03) 0.1 (0.00)</td>
</tr>
<tr>
<td>Bladder</td>
<td>0.5 (0.01) 0.0 (0.00) 0.5 (0.01) 0.3 (0.00)</td>
</tr>
<tr>
<td>Uterus</td>
<td>0.8 (0.01) 0.3 (0.00) 1.2 (0.01) 0.4 (0.00)</td>
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<tr>
<td>Ovaries</td>
<td>0.5 (0.01) 0.0 (0.00) 0.6 (0.01) 0.1 (0.00)</td>
</tr>
<tr>
<td>Vagina</td>
<td>0.5 (0.01) 0.0 (0.00) 0.3 (0.00) 0.1 (0.00)</td>
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* Picomole value of specific tissue divided by the sum of picomole values of all measured tissues × 100. n 3 per time point.
production of lignan metabolites, intestinal absorption or serum kinetics is an unlikely explanation, as no differences in serum, urine or faecal concentrations were observed between males and females. It is possible, however, that the pharmacokinetics of lignans differs between sexes in these particular tissues, and may result in diverse treatment responses between sexes. Thus, identification of those tissues with a sex difference in lignan concentrations is important when assessing their effectiveness as well as safety.

In the present study, the calculated contribution of tissue residual blood to tissue radioactivities was low, did not differ significantly between sexes and duration of exposure and did not explain the observed differences between sexes or tissue distributions.

In conclusion, prolonged \(^3\)H-SDG exposure increased radioactivity excretion and serum and tissue concentrations in both males and females. The significant access of SDG metabolites into tissues indicates the possibility for local effects \textit{in situ}. In specific tissues, accumulation still occurred after 7d exposure regimen when serum concentrations were already stabilised, indicating that serum concentrations alone do not fully reflect specific tissue concentrations. Moreover, observed sex differences in lignan tissue distribution indicate the possibility for distinct treatment responses in males and females. These findings will facilitate the identification of target tissues of lignan actions, and will help in the better understanding of the mechanisms of lignan action \textit{in vivo}.

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