

# Annex A - Information on ginger

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8. Ginger (**Zingiber officinale**) is a flowering tropical plant originating in Southeast Asia and grown in warm climates including China, India, Africa and the Caribbean. Ginger is commonly consumed in fresh root, dried root powder and capsule (encapsulated dried powder) forms, as a liquid extract, preserved in syrup or sugar and as a tea.

## Uses

9. The rhizome (underground stem) of the ginger plant is commonly used as a spice and flavouring in many countries around the world and is increasingly growing in popularity as a natural remedy due to its purported immune system-boosting properties and also to alleviate motion sickness and post-operative nausea and vomiting. Ginger has been recommended for the treatment of pregnancy-related nausea (NHS 2021; NICE 2021) and has also been used as a dietary supplement and a traditional remedy in many cultures. Ginger is included in the official pharmacopoeias of some western countries.

## Constituents

10. Over 100 compounds have been identified in ginger extracts, the main being terpenoids - mainly sesquiterpenoids ( $\alpha$ -zingiberene,  $\beta$ -sesquiphellandrene,  $\beta$ -bisabolene,  $\alpha$ -farnesene, ar-curcumene (zingiberol) and smaller amounts of monoterpenoids (camphene,  $\beta$ -phellandrene, cineole, geraniol, curcumene, citral, terpineol, borneol) (EMA, 2012).

11. The ginger rhizome contains two main classes of constituents: the essential oils responsible for the aroma, and the main pungent principles, gingerols and shogaols. Organic acids are also present in smaller amounts. Depending on the area of cultivation, gingerols make up 4-7.5% of the pungent principles, the main one being 6-gingerol. Gingerols of other chain lengths are also present in smaller amounts.

## **Reviews by other regulatory agencies**

12. Ginger is included in the official pharmacopoeias of several western countries. Ginger is classified as 'Generally Recognised as Safe' (GRAS) by the United States Food and Drug Administration (FDA) however, few specific studies have been carried out to evaluate the safety of ginger use during pregnancy and lactation. A report by the National Institute for Health and Care Excellence (NICE) cites a number of short duration trials which have been conducted in pregnant women (NICE, 2021).

13. In 2008, the Danish company Ferrosan A/S withdrew their product GraviFrisk - a product containing 6 g of dried ground ginger - from market, due to concerns surrounding the lack of safety data with respect to the use of supplements containing highly concentrated ginger extracts by pregnant women (Dietz *et al.*, 2016).

14. In their 2012 report on ginger root in powdered form, the European Medicines Agency (EMA) concluded "The ginger extract dosages to provoke acute toxicity are high and much higher than usually administered dosages (factor 10-15 for an adult). There is some evidence that ginger root may cause testicular weight to increase by repeated high dosages of ginger root extract (2000 mg/kg). Ginger root has mutagenic as well as antimutagenic properties in microbial test systems. Developmental toxicity studies in rats are difficult to interpret, however, it is probably not a cause for concern. In general, toxicity studies of ginger are considered inadequate at least regarding genotoxicity, carcinogenicity and, partially, reproductive and developmental toxicity."

15. The Norwegian Food Safety Authority have issued a warning regarding the use of ginger supplements and ginger-containing shots during pregnancy. This was based on a risk assessment carried out by the Danish Technical University and the Danish Veterinary and Food Administration (DTU, 2018). The assessment, based on animal studies, including one where rats were treated with a fresh grated ginger preparation with ginger at concentrations of 20-50 g/L in water, found that even in the 20 g/L treatment group – the equivalent of 1,784 mg/kg bw increased the incidence of abortion in rats. The Norwegian Food Safety Authority concluded that while a woman of 70 kg would consume less ginger (124 mg to 329 mg) there remains cause for concern and fetal risk cannot be excluded.

16. Recently, the Finnish Food Authority has issued a recommendation against the use of products containing ginger concentrate or extract, ginger tea and food supplements containing ginger by pregnant and breastfeeding women, infants and toddlers, schoolchildren, elderly and individuals with weakened immunity (Finnish Food Authority, 2019). It was noted that the concentrates contained harmful substances and safe consumption levels were unknown.

## **Health-based guidance values**

17. There are currently no health-based guidance values (HBGV) with respect to ginger or its main components. Exposure to ginger was considered based on information found on supplement and tincture composition and background diet, but the variability of available supplements meant exposure also varied, which made comparison difficult.

18. The NHS and NICE recommend the use of ginger tea and biscuits for the treatment of nausea and vomiting in pregnancy (NHS, 2021; NICE, 2021). Generally, amounts of 1 g ginger per day have been recommended (NHS, 2022). Anecdotally, 1-1.5 g per day has also been advised during pregnancy (Healthline, 2020; Mother and baby, 2022). The use of supplements under the advice and supervision of a medical professional is advised.

## **Toxicology**

19. It was noted that some studies detailed effects on male testes and, though not relevant for females, they were nevertheless regarded as indicating a potential reprotoxic effect from ginger. Studies suggest that ginger affected the viability of pregnancy; however, with no strong conclusive human data, the COT

concluded that more work was required, especially as these studies suggested a link between first trimester loss and ginger use. Further, the possible fetotoxicity based on evidence from animal data, genotoxicity and possible drug interactions should be further investigated.

20. Discussion paper TOX/2021/26 reviewed the available studies on cytotoxicity, mutagenicity, acute, reproductive and developmental toxicity, lactation and possible drug interactions as well as data on potential exposure in pregnancy, covering both animal and human studies. The results from these reports were varied due to the differences in the forms and extracts tested and as a result, some findings were contradictory.

21. Paper TOX/2021/51 provided further information with respect to animal studies, contaminants and exposure to ginger supplements, primarily centred on the effect of ginger on prostaglandins, reproductive and developmental toxicity and the possible contaminants present in ginger. The Committee noted that the papers reviewed covered ginger in a range of forms including fresh, aqueous, dried and alcohol extracts.

## **Cytotoxicity**

22. The cytotoxicity of ginger has been investigated with varied results. Plengsuriyakarn *et al.* (2012) examined cytotoxicity in Cholangiocarcinoma (CCA) cell line 6 (CL-6), hepatocarcinoma (HepG2) and normal human renal epithelium (HRE) models using calcein-AM release and Hoechst 33342 assays. The median inhibitory concentration, (IC<sub>50</sub>) and value for cytotoxicity of the crude ethanolic extract of ginger were 10.95 and 53.15, µg/ml respectively.

23. Zaeoung *et al.* (2005) reported that the IC<sub>50</sub> of ginger was higher than 39.2 µg/ml against breast (MCF7) and colon (LS174T) cell lines. Abudayyak *et al.* (2015) found the aqueous and methanol extracts of ginger exhibited no cytotoxic activity when assessed using an MTT test (a colourimetric assay for assessing cell metabolic assay) in the rat kidney, NRK-52E cell line. The chloroform extract resulted in an IC<sub>50</sub> value of 9.08 mg/mL.

24. However, it was noted that the inhibitory concentration (IC<sub>50</sub>) values presented in the studies reviewed were based on a small amount of data, from only six different cell lines and therefore firm conclusions could not be drawn. Also, the purpose of most of these studies was an attempt to identify possible anti-cancer agents, rather than as an assessment on the safety of ginger as a supplement and therefore relevant endpoints would not have been assessed.

## Mutagenicity

25. Nakamura and Yamamoto (1982) found that the juice of ginger rhizome possessed both mutagenic and anti-mutagenic properties, and that 6-gingerol in particular was a powerful mutagen. The group also demonstrated that 6-shogaol was much less mutagenic (strain Hs30 of *Escherichia coli*) than 6-gingerol (Nakamura & Yamamoto 1983). In a ***Salmonella typhimurium*** reverse mutation (Ames) assay, the urine of rats fed diets containing 0.5, 1 and 5% powdered ginger for 1 month and exposed to benzo(a)pyrene was found to display a significant reduction in the mutagenicity as indicated by a reduced number of TA98 and TA100 revertants at all ginger concentrations (Nirmala *et al.* 2007) when tested in an Ames assay.

26. In another Ames assay, an ethanol extract of ginger (Soudamini *et al.* 1995) and an essential oil from ginger (Sivaswami *et al.* 1991) demonstrated mutagenic activity in *S. typhimurium* strains TA 100 and TA 1535 at concentrations of 25-50 mg/plate and 5-10 mg/plate, respectively. Similarly, an ethanolic ginger extract at concentrations between 10 and 200 µg/plate, and gingerol and shogaol were mutagenic in strains TA 100 and TA 1838 with metabolic activation by rat liver S9 fraction, while zingerone did not display mutagenic effects (Nagabhushan *et al.* 1987).

27. Abudayyak *et al.* (2015) found the aqueous ginger extract exhibited mutagenic activity when assessed using the Ames assay on *S. typhimurium* TA98 (-S9) strains. No activity was observed with the chloroform and methanol extracts.

28. Based on the available data, ginger showed some mutagenicity in TA 100, TA 1535, and TA 98 strains, but this is low compared with established mutagens. Ginger is not shown to be mutagenic in vivo (Nirmala, Prasanna Krishna and Polasa, 2007).

## Acute toxicity

29. An acute toxicity study carried out by Malik and Sharma on male Wistar rats showed no signs of toxicity or mortality (2011). The animals were administered doses of 250, 500 and 1000 mg/kg lyophilised ginger powder by gastric gavage. The authors stated that the three dose levels used in the study corresponded to 5, 10 and 20% of the NOAEL of the powder (5000 mg/kg).

30. Plengsuriyakarn et al. also examined acute and subacute toxicity in hamsters. Sixty hamsters (nine groups of 5 male and 5 female) were fed either 1000, 3000 or 5000 mg/kg bw ethanolic ginger extract, resuspended in a distilled water-Tween-80, mixture 4:1, v:v, by oral gavage and observed for 14 days (acute toxicity) or 30 days (subacute toxicity). The group concluded that there was an absence of any toxicity at maximum dose of 5 g/kg bw during the investigation period.

### **Short term repeat dose studies**

31. Rong et al. (2009) evaluated the safety of powdered Japanese ginger (mainly containing 6-gingerol galanolactone and 6-shogaol) by conducting a 35-day toxicity study in rats. Both male and female rats were treated with 500, 1000 and 2000 mg/kg bw/day by gavage. The results demonstrated that oral administration of up to 2000 mg/kg to male and female rats did not result in any increase in mortality, or changes to behaviour, growth, the general condition of the animals (including: changes in skin, fur, eyes, and mucous membranes, occurrence of secretions, excretions and autonomic activity), food and water consumption. It was only at the highest dose tested (2000 mg/kg), that ginger led to slightly reduced absolute and relative weights of testes (by 14.4% and 11.5%, respectively). No effects were apparent in the females.

32. The effect of oral and intraperitoneal administration of aqueous extracts of ginger root over 28 days in female rats at two dose levels (50 mg/kg and 500 mg/kg) was examined for haematological, serum and systemic toxicity (Alnaqeeb et al. 2003). Neither oral nor intraperitoneal administration resulted in mortality. Orally administered aqueous ginger extract resulted in increased levels of serum aspartate aminotransferase (AST) and decreased levels of alanine aminotransferase (ALT).

33. Jeena et al., conducted a sub chronic toxicity study of the essential oil of ginger in Wistar rats following oral administration at concentrations of 100, 250, and 500 mg/kg per day once daily for 13 consecutive weeks to assess the oral safety of ginger oil (2011). No mortality was observed. No unusual changes in behaviour or locomotor activity were observed during the period of the study, nor were any abnormal changes observed in the relative organ weights of liver, kidney, spleen, lungs, brain, and stomach with respect to body weight in ginger oil-treated animals when compared to vehicle control animals.

34. An increase in serum sodium levels was observed in male rats treated with 500 mg/kg per day but in the absence of changes in sodium levels in females, this change was not considered significant. A slight increase in total bilirubin was observed in female rats treated with ginger oil along with a decrease in AST and ALT levels however, there was no significant changes in hepatic function parameters such as alkaline phosphatase, total protein, albumin, and globulin content.

## **Reproductive and developmental toxicity**

### **In vitro studies**

35. Mohammed *et al* investigated the effects of herbal extracts, including ginger and 6-gingerol, on chick embryonic heart micromass and mouse D3 embryonic stem cell systems (ESD3) (2016). The team observed that the use of ginger herbal remedies in the first trimester of pregnancy may affect foetal development. However, 6-gingerol-treated primary embryonic chick cardiomyocytes showed no significant changes in contractile and cellular activity or changes in total protein content in comparison to the control.

36. At concentrations of 0.75–6  $\mu\text{M}$ , 6-gingerol treated primary embryonic chick cardiomyocytes exhibited no significant changes in contractile activity, cellular activity or changes in total protein content in comparison to the control. At concentrations of 12.5–50  $\mu\text{M}$ , inhibition in contractile activity was observed at 48h. All high 6-gingerol concentrations, 12.5–100  $\mu\text{M}$ , tested in micromass, significantly altered both the cellular activity and protein content in a dose-dependent manner.

37. The same concentrations of 6-gingerol were used to treat the ESD3, which showed a significant decrease in cardiomyocyte differentiation for all tested concentrations except 0.75  $\mu\text{M}$ . The cellular activity and protein content of stem cell-derived cardiomyocytes also exhibited a significant decrease with increased 6-gingerol concentration exposure. The 6  $\mu\text{M}$  dose showed almost no differentiation in the morphological score on days 10 and 11, which was considered an anomaly. However, a small amount of differentiation was observed on day 12.

### **Animal studies**

38. To date, the number of studies on the safety of the use of ginger supplements during pregnancy is limited. The ginger component 6-gingerol, was

highlighted to affect some essential embryonic developmental processes, such as the disruption of angiogenesis. Kim *et al*, demonstrated the ability of 6-gingerol to inhibit proliferation and tube formation of primary cultured human endothelial cells in rat aorta by down regulation of cyclidin D and the ability to inhibit tumour growth in mice through its anti-angiogenic activity (2005).

39. The animal studies on reproductive toxicity considered in TOX/2021/26 reported a number of findings, including reduced maternal weight gain, increased fetus weight, increased serum testosterone level in F1 generation males and an increase in embryonic loss.

40. The teratogenicity of EV.EXT 33, a patented Zingiber officinale extract (comprising 6-gingerol, 8-gingerol, 10-gingerol, 6-shogaol, and 8-shogaol, which made up 1.9 w/w of the extract) was investigated in Wistar rats, (Weidner & Sigwart, 2001). The extracts were administered orally by gastric intubation at concentrations of 100, 333 and 1000 mg/kg, to three groups of pregnant rats from days 6 to 15 of gestation. Their bodyweight, food and water monitored during the treatment period. The study concluded that treatment with EV.EXT 33 during the period of organogenesis resulted in neither maternal nor developmental toxicity at daily doses of up to 1000 mg/kg bw.

41. Shalaby and Hamowieh investigated fertility, serum testosterone and acute toxicity of ginger in rats (2010). One hundred and twenty male Sprague Dawley rats, separated into groups of 10, were orally administered either water (prepared using 100 g dry ginger roots soaked in 500 ml water or 500 ml methyl alcohol 90%) or methanolic extracts in graded doses ranging from 5 to 17.5 g/kg bw (gavage doses were not specified). Following dosing, the number of dead mice in each group after 48 hours of observation were recorded. The oral Lethal Doses (LD50) of the methanolic and water extracts were calculated to be 10.25 and 11.75 g/kg bw respectively. No symptoms of toxicity were observed at does up to 5 g/kg bw. Both extracts increased fertility index, sexual organ weight, and sperm motility and count after 65 consecutive days (see below).

42. To investigate the effect of ginger extracts on serum testosterone levels, male rats had their fertility reduced by inducing diabetes, a condition shown to reduce male fertility. The aim was to see whether ginger, with its antioxidant and androgenic effects, would restore fertility. Rats were rendered diabetic by subcutaneous injection of 120 mg/kg bw alloxan for 3 days, were administered methanolic extracts of ginger for 65 days at doses of 100 and 200 mg/kg bw/d. Testosterone levels increased to  $4.08 \pm 0.10$  and  $7.13 \pm 0.14$  ng/dL (both significant at  $P 0.001$ ) compared to the diabetic control group who had



levels of  $3.30 \pm 0.03$  ng/dL. Serum testosterone levels also increased in rats given water extracts (150 and 300 mg/kg bw) were determined to be  $4.06 \pm 0.03$  and  $5.04 \pm 0.08$  ng/dL (both significant at  $P 0.001$  when compared to the diabetic control group) respectively.

43. The team also investigated fertility with regards to fertility index (for each male this was calculated as percentage of number of females that become pregnant in relation to number of mated females) and spermatogenesis. Rats were orally administered methanolic extracts at doses of 100 and 200 mg/kg bw for 65 consecutive days and water extracts at doses of 150 and 300 mg/kg bw and compared to a diabetic control group.

44. Histopathological examination of the testes of diabetic rats showed mild to moderate degenerative changes of spermatogenic cells, diffuse oedema and incomplete arrest of spermatogenesis. The testes of rats orally administered 300 mg/kg bw of water extract of ginger roots showed mild degeneration of spermatogenic cells and slight oedema of interstitial cells. The testes of rats receiving orally 200 mg/kg bw of methanolic extract of ginger roots showed nearly normal seminiferous tubules, showing fewer signs of degradation, suggesting a LOAEL of 200 mg/kg bw/day for the methanolic extract. The team concluded that the results suggest the intake of ginger root as a drink may be useful for diabetic patients suffering from sexual impotency.

45. The above study has been included for completeness and as any general mechanisms may be more widely relevant: This is consistent with the findings of Hosseini et al (2015) - see paragraph 29, Annex B.

46. No additional reproductive studies were identified. Therefore, any point of departure for ginger would need to be based on the animal studies discussed in TOX/2021/26 and reproduced in paragraphs below.

47. Reproductive and developmental toxicity has also been investigated in rat studies. In a study by Wilkinson (2000), three groups of pregnant Sprague-Dawley rats were administered either a control (unspecified), or 20 g/L or 50 g/L ginger tea - prepared by the infusion of grated ginger in water via their drinking water - during days 6 to 15. No further details were provided regarding specific compounds of interest. While no maternal toxicity was observed, embryonic loss in the treated groups was found to be double that of the controls. Exposed foetuses were found to be significantly heavier than controls and showed no gross structural malformations. The results of this study suggest that in utero exposure to ginger tea results in early embryonic loss and increased growth in surviving

foetuses.

48. Hosseini et al. investigated the effect of ethanolic ginger extract on serum testosterone, LH and FSH as well effect on spermatogenic cell lines in male mature offspring rats (2015, abstract only). In this study, 72 female rats, sorted into 9 groups were orally administered of alcoholic extract of ginger at doses of 50, 100 and 200 mg/kg bw, during their neonatal and perinatal periods versus saline as a control. Following puberty, LH, FSH, cell numbers of Sertoli, spermatogonia, spermatocyte and spermatids were counted in 8 male rat offspring from each group. Ginger was found to significantly increase testosterone levels and the number of spermatogenic cells and at doses of 100 and 200 mg/kg bw, alcoholic extract of Ginger significantly reduced the FSH and LH compared to control groups. The authors concluded that “the oral consumption of Ginger during pregnancy and lactation dose-dependently increase the level of testosterone and the number of spermatogenic cells.”

49. The teratogenicity of EV.EXT 33, a patented *Zingiber officinale* extract (comprising 6-gingerol, 8-gingerol, 10-gingerol, 6-shogaol, and 8-shogaol, which made up 1.9 w/w of the extract) was investigated in Wistar rats, (Weidner & Sigwart, 2001). The extracts were administered orally by gastric intubation at concentrations of 100, 333 and 1000 mg/kg, to three groups of pregnant rats from days 6 to 15 of gestation. Their bodyweight, food and water monitored during the treatment period. The study concluded that treatment with EV.EXT 33 during the period of organogenesis resulted in neither maternal nor developmental toxicity at daily doses of up to 1000 mg/kg bw.

50. Dissabandara & Chandrasekara (2007) also examined the effect of powdered ginger extract administered prenatally on the postnatal development of rats. A period of administration of the dry powdered extract orally at doses of 500 mg/kg/day or 1000 mg/kg/day (control not specified) during days 5 to 15 of gestation resulted in a lower intake of food and water and lower weight gain in the ginger treated group, suggesting that maternal administration of ginger during mid pregnancy resulted in reduced maternal weight gain and increased embryonic loss without affecting the postnatal growth and physical maturation of the surviving offspring.

51. ElMazoudy and Attia (2018) investigated the effect of powdered dried ginger root on the oestrus cycle and implantation in female mice. ICR mice, orally dosed at 250, 500, 1000, or 2000 mg/kg bw/d aqueous ginger extract. These were investigated in four different experiments: the main study of outcomes (treatment for 90 days and throughout mating and gestation), a 35-day treatment study

evaluating effects on the oestrous cycle. The third and fourth intended antifertility and abortifacient loss (20 days treatment). In the main study, the dams were sacrificed on gestation day 20. One mortality was recorded in the 100mg/kg bw/d group on gestation day 18 and two mortalities in the 2000 mg/kg bw/d group at day 16. There was also a significant reduction in body weight change in these two dose groups compared to the control group; however, food consumption was comparable.

52. In the study investigating the oestrus cycle, a significant reduction in the numbers of oestrus cycles was observed at the highest dose group, with the length of the oestrus cycle in this group being significantly prolonged ( $10.05 \pm 0.8$ ) days compared with ( $4.99 \pm 0.5$ ) days recurrent and successive oestrous cycles in control mice. At the highest dose level, the length of the oestrous cycle was prolonged with a significant decrease in the duration of diestrus-metestrus (luteal) phase and prolonged proestrus-estrus (ovulatory) phase. In the study investigating pre-implantation loss, a significant decrease in the number of corpora lutea was observed at the highest dose group. Implantation failure was also increased by 36% compared to the control group and pre-implantation loss at this dose group was also 16.59% higher than the control group. The authors considered that this may reflect a dose-dependent antifertility (anti-implantation) effect.

53. Regarding fertility and developmental outcomes, the female copulation index was significantly reduced at 2000 and 1000 mg/kg bw/d groups, whereas the female pregnancy index was significantly decreased only at the highest dose. The number of implantation sites and live fetuses in the 2000 mg/kg bw/d group was lower than the other treated and control groups. An increase in fetal resorption and post implantation loss was also seen at the highest dose group. There was no evidence of fetal malformations however growth retardation, reduced pup weight and delay in the crown-rump length was observed at this dose group as well. Finally, changes in ovarian histopathology were observed at 2000 mg/kg bw/d, following 90 days of treatment. Ovarian follicle atresia was observed. The atretic follicles contained cell debris and there was haemorrhage in the antral cavity.

54. Additionally, degenerated primordial follicles with pyknotic nuclei forming polycystic ovaries were noted. Deteriorated follicles were observed as a detaching of layers of granulosa cells from the basal membrane by dilation of zona pellucida and with evidence of apoptosis in damaged ova follicular nuclei were non-visible. The authors considered the above observations as evidence that

ginger possesses anti-ovulation properties. Overall, the authors concluded that ginger impairs the normal growth of corpus luteum because of progesterone insufficiency during early pregnancy and that the results suggested that ginger can disrupt the oestrous cycle and blastocyst implantation without teratogenesis. They considered the highest NOAEL to be 500 mg/kg bw.

55. The Committee considered the animal studies to be inconclusive. Plengsuriyakarn *et al.*, examined acute and subacute toxicity in hamsters (2012). Sixty hamsters (nine groups of 5 male and 5 female) were fed either 1000, 3000 or 5000 mg/kg bw ethanolic ginger extract, resuspended in a distilled water-Tween-80, mixture 4:1, v:v, by oral gavage and observed for 14 days (acute toxicity) or 30 days (subacute toxicity). The group concluded an absence of any toxicity at maximum dose of 5 g/kg bw during the investigation period.

56. ElMazoudy and Attia (2018) noted reductions in bodyweight and mortalities in mice dosed up to 2000 mg/kg bw/day and Alnaqeeb *et al.*, (2003), observed increases in serum aspartate aminotransferase (AST) in female rats dosed up to 500 mg/kg. However, the Committee noted that the database was limited and on the basis of the available information, more data would be needed in order to allow for a robust investigation of the effects described above. Therefore, at present, the Committee were unable to determine a point of departure, to reach a conclusion.

### **Human studies**

57. The COT considered a number of epidemiological studies investigating the use of ginger during pregnancy ([TOX/2021/26](#)). For the most part, few studies explicitly addressed the safety of ginger consumption during pregnancy. Most were focused on the use of ginger as a treatment for nausea (Fischer-Rasmussen *et al.*, 1990; Smith *et al.*, 2004; Ensiyeh *et al.*, 2009), age-related neurological disorders or pregnancy-induced sickness and therefore focused on efficacy (Willetts *et al.*, 2003; Stanisiere *et al.*, 2018). However, safety was considered in a few studies. The studies considered by the Committee included observational and randomised clinical studies, lasting from 4 days to 20 weeks in duration (Vutyavanich *et al.*, 2001; Portnoi *et al.*, 2003). Ginger in various forms was investigated in doses ranging from 750 mg/day to the equivalent of 7 g/day.

58. The study results in pregnant women were also varied and the overall findings inconclusive. Findings reported included abdominal discomfort, vomiting and diarrhoea. There were reports of incidences of spontaneous abortion (Portnoi *et al.*, 2003, Ensiyeh *et al.*, 2009), however, this effect was observed in both the

treated and control groups and therefore, cannot directly be attributed to the consumption of ginger. Portnoi *et al.*, reported 8 spontaneous abortions in the comparator group, compared to 3 occurring in the group taking ginger and Ensiyeh *et al.*, reported 2 spontaneous abortions in the ginger group compared to 1 in the group taking vitamin B6. This study reported no effects of defects post-partum following exposure to ginger.

### **Human studies - exposures in pregnancy**

59. In their 2003 review of interventions for nausea and vomiting in early pregnancy, Jewell and Young concluded that maternal consumption of ginger shows no evidence of teratogenicity in infants (Jewell and Young, 2003). More recently, Stanisiere *et al.* (2018) conducted a review of the safety and efficacy of ginger rhizome for decreasing nausea and vomiting in women during early pregnancy. This review was based on systematic literature searches until the end of December 2017. Most of the studies included in this review have already been included in this draft statement. Some recent studies have been conducted evaluating the effectiveness and safety of ginger in pregnancy, and these will be discussed in detail. Overall, most studies reported gastrointestinal effects such as abdominal discomfort, vomiting and diarrhoea. Other effects included dizziness, headaches and drowsiness with some more serious effects such as spontaneous abortion also being reported in 5 out of the 14 randomized clinical studies. The review by Jewell and Young focuses on the reported effects rather than statistical significance, therefore more details on the studies reporting more serious effects are given below.

60. In a double-blind randomised crossover trial, 27 women were administered capsules containing either 250 mg ginger in powdered root form or 250 mg lactose as a placebo, four times per day, for four days followed by a wash out period of 2 days prior to a further 4 days administration of ginger or placebo (Fischer-Rasmussen *et al.*, 1990). Two subjects did not carry to term: One subject from the ginger group had a spontaneous abortion, one elected. Of the remaining 25 subjects, no adverse effects were observed.

61. Of the available human studies, few explicitly addressed the safety of ginger consumption during pregnancy, most being incidental to other studies. In a double-blind study by Vutyavanich *et al.* (2001), 32 women were given 1 g of dried ginger in capsule form for 4 days. Of those in the ginger group, one spontaneous abortion was reported compared to 3 in the placebo group. Equally, for delivery by caesarean section, there was no difference between both groups.

No congenital abnormalities were observed in all babies carried to term. The group concluded that there were no significant adverse effects of ginger on pregnancy outcome.

62. An observational study in humans examined 187 pregnant women who took ginger in their first trimester and compared them to 187 pregnant women exposed to nonteratogenic drugs that were not antiemetic drugs. The results suggested that the ginger group did not have an increased rate of major malformations above the baseline rate of 1%–3% (Portnoi et al., 2003). Three major malformations were reported in the ginger group, ventricular septal defect (VSD), right lung abnormality, and kidney abnormality (pelviectasis) and one child was diagnosed with idiopathic central precocious puberty at age 2 years. The mother was reported to have taken 250 mg of ginger in capsules four times a day from 11 to 20 weeks of gestation in addition to dimenhydrinate and doxylamine/vitamin B6 (Diclectin) during the first trimester of pregnancy. No significant difference between the two groups in terms of live births, spontaneous abortions, stillbirths, therapeutic abortions, birth weight, or gestational age were reported, however the comparison group had more infants weighing less than 2500 g and the ginger group had 8 sets of twins.

63. Willetts et al. examined the effect of ginger on pregnancy induced nausea (2003). 120 women less than 20 weeks pregnant, were given 125 mg ginger extract (EV.EXT35; equivalent to 1.5 g of dried ginger) or a placebo four times per day for 4 days. Three spontaneous abortions were observed in the group receiving ginger compared to those in the placebo group. Whilst also examining the use of ginger in the treatment of nausea and vomiting in pregnancy, Smith et al. noted 3 spontaneous abortions in the group taking 1.05 g ginger compared to 9 in the group taking 75 mg B6 daily for 3 weeks (2004).

64. Ensiyeh et al, investigated the effectiveness of ginger versus B6 for treatment of NVP (2009) in women before 17 weeks' gestation. 70 women were randomised to receive either ginger at a dose of 1 g per day or B6 and 40 mg per day for 4 days. The ginger group reported 2 spontaneous abortions, compared to one in the B6 group. Of the babies brought to term, no congenital anomalies were observed, and all babies were discharged in good condition.

65. The COT considered the mode of action of the purported benefits of ginger on nausea. It was theorised that ginger might decrease prostaglandin levels, which were linked to nausea.

66. The Committee noted the potential effect of ginger on the prostaglandin pathway, in particular Cyclooxygenase-1 (COX1) and Cyclooxygenase-2 (COX2) inhibition and how this may affect early pregnancy. One study examining the effects of ginger extracts on prostaglandin E2 (PGE2) production *in vitro* (Lantz *et al.* 2007) demonstrated that crude organic extracts (dichloromethane-methanol, 1:1 v/v) of ginger were capable of inhibiting PGE2 production and that the compounds may act at several sites. The most potent effect on lipopolysaccharide (LPS) induced prostaglandin production was noted at less than 0.1 µg/ml. It was noted that the half maximal inhibitory concentration (IC50) values for a range of components were given, and it was demonstrated that the components mainly acted on COX-2. The COT concluded further studies would be needed to determine the role of increased prostaglandin levels in the early termination of pregnancy.

67. Overall, it was concluded that there were limited data. The human data presented were not strongly indicative of any toxicological concern but there were some indications of possible effects and many uncertainties. Ginger did not appear to be systemically toxic but reprotoxic effects have been reported in animal studies. However, there is no significant evidence for this outcome in human studies.

## **Lactation**

68. With respect to lactation, the focus of available studies (**Lamxay *et al.*, 2011; Kaygusuz *et al.*, 2021; Dilokthornsakul *et al.*, 2021**) has been on the effect of ginger on milk production and volume rather than safety and therefore, the effect of exposure during lactation has not been fully investigated.

## **Effect on Cytochrome P450 (CYP) Enzymes and Herb-Drug Interactions**

69. CYPs are a family of enzymes responsible for the biotransformation of several drugs. Induction or inhibition of CYP enzymes is a major determinant of the occurrence of drug-drug interactions.

### **In silico**

70. Qiu *et al.* (2015) investigated the molecular interactions between 12 main active components (6-gingerol, 8-gingerol, 10-gingerol, 6-shogaol, 8-shogaol, 10-shogaol, ar-curcumene, β-bisabolene, β-sesquiphelandrene, 6-gingerdione, (-)-zingiberene, and methyl-6-isogingerol) and human cytochrome P450 (CYP) 1A2, 2C9, 2C19, 2D6, and 3A4 and attempted to predict the

absorption, distribution, metabolism, excretion, and toxicity (ADMET) of the 12 ginger components using computational methods and literature searches. This study suggests that ginger components may regulate the activity and expression of various human CYPs, resulting in alterations in drug clearance and response.

71. These results could potentially be significant in pregnant women on medication, who are using ginger as a remedy for nausea in the early stages of pregnancy.

#### **In vitro studies**

72. Ginger extracts and the major components thereof - 6-gingerol (6G), 8-gingerol (8G), 10-gingerol (10G) and 6-shogaol (6S) - were investigated in *in vitro* models and shown to have an inhibitory effect on CYP isoenzymes CYP3A4, CYP2C9 (Kimura *et al.*, 2010) CYP2C19 (Kim *et al.*, 2012) and CYP1A2 and CYP2C8 with IC50 values as low 1 $\mu$ M, (e.g., 6-shogaol on CYP1A2; Mukkavilli *et al.*, 2014).

#### **Animal studies**

73. Several reports have been published on the pharmacological properties of ginger, with varying results. Studies have examined herb-drug interaction in animal models, (Okonta *et al.*, 2008; Egashira *et al.*, 2012) although some studies have questionable results.

74. A study into the effect of ginger on the pharmacokinetics of metronidazole was studied by Okonta *et al.*, using rabbits (2008). In a two-phase study, five healthy local strain rabbits (3 females, two males) were each given 3 mg/kg oral metronidazole. Following a 2-week washout period, the rabbits were given 1 ml/kg of ginger extract orally daily for 3 days and immediately given 3 mg/kg metronidazole per oral on the third day. Ginger significantly increased the absorption and plasma half-life, and significantly decreased the elimination rate constant and clearance of metronidazole.

75. Egashira *et al.*, reported the interaction between ginger juice and tacrolimus in rats (2012). Tacrolimus (0.6 mg/kg) was administered intraduodenally in male Sprague-Dawley rats 1 hr following oral administration of 10 mL/kg 50% ginger juice or water. Pre-treatment with ginger juice was found to significantly increase tacrolimus blood concentrations compared to those pre-treated with water or orange juice.



76. The possible herb-drug interaction of ginger crude extract on glibenclamide and insulin was investigated by Al Omari *et al.*, along with its hypoglycaemic and antihyperglycemic effects in normoglycemic- and streptozotocin-induced (STZ) diabetic rats (2012). Ginger crude extract was administered to normoglycemic male rats as a single dose (1 day) and as a daily dose for 1 week. STZ diabetic rats were treated with the same GCE concentrations (25, 50 and 100 mg/kg bw) together with glibenclamide (5 mg/kg bw) or insulin (1.2 IU/kg bw).

77. Single administration of ginger crude extract showed a significant decrease in blood glucose level (BGL) in normoglycemic rats after 1 and 2 hours (50 mg/kg bw). In STZ- diabetic rats ginger crude extract (25 and 50 mg/kg bw) decreased non-fasting BGL (N-FBGL) significantly at 1.5, 2.5, 3.5 and 4.5 hours. Glibenclamide (5 mg/kg bw) in combination with ginger crude extract at doses 25 or 50 mg/kg bw exhibited a significantly reduction in the N-FBGL 26.3% and 25.1% respectively after 4.5 hours, compared to glibenclamide alone which exhibited a 7.9% reduction.

#### **Human studies**

78. Human data showed possible interactions with medicines, including antibiotics, immunosuppressants, and anticoagulant medications. Although, in some cases, multiple concomitant medications were being used therefore, the effects observed cannot necessarily be directly attributed to ginger supplementation (Rubin *et al.*, 2019).

79. Conversely, whilst investigating the effects of ginger on the pharmacokinetics or pharmacodynamics of warfarin and the effect of ginger on clotting status, Jiang *et al.*, (2005), found that neither the pharmacokinetics nor pharmacodynamics of warfarin were affected in healthy males who were treated with a single 25 mg dose of warfarin, following 7 days of pretreatment with ginger tablets (3 tablets, 3 times per day, each capsule containing extract equivalent to 0.4 g of ginger rhizome powder). Furthermore, ginger had no effect on international normalized ratio (INR) or *ex vivo* platelet aggregation in response to arachidonic acid.

80. Ginger was found to have a significant inhibitory effect on CYP3A4, CYP2C9, and P-glycoprotein activities *in vitro* (Kimura *et al.*, 2010; Zhang and Lim, 2008). It was this effect that was thought to be responsible for reported hepatic cytolysis in a 48-year-old woman being treated with crizotinib. The patient, who was being treated with 250 mg crizotinib twice a day, had been taking ginger as a

tea (amounts unknown) concomitantly during treatment. A subsequent diagnostic evaluation showed an increased crizotinib concentration, 1.8-fold higher than that measured two months prior.

### **Anti-platelet aggregation activity**

81. Several reports have been published on the pharmacological properties of ginger, with varying results. The potential effect of ginger extract and components thereof on the reduction of platelet aggregation and their potential antithrombotic activity has been noted as a concern in both literature and by health professionals.

82. Ginger was reported to have antiplatelet activity (Srivastava, 1986,1989; Young *et al.*, 2006), with some studies reporting effects in animals at doses of 500 mg/kg bw (Thomson *et al.*, 2002). Ginger was found to inhibit platelet thromboxane and prostaglandin endoperoxidases (PGF<sub>2</sub> $\alpha$ , PGE<sub>2</sub> and PGD<sub>2</sub>) in human platelets, in a dose-dependent manner (Srivastava, 1984).

83. With regards to the relevance of such effects in pregnancy, literature reports note that pregnancy is associated with an increased incidence of thrombotic events; mainly related to a pro-thrombotic state, physiologically useful to reduce bleeding at delivery. These changes are more pronounced in the third trimester (Patti *et al.*, 2014). It has also been hypothesised that antiplatelet agents might prevent or delay the development of pre-eclampsia (Duley *et al.*, 2019). The implications and clinical significance of the anti-platelet activity of ginger exposure during different stages of pregnancy remain undetermined.

84. This further highlighted the need to differentiate exposure from the normal diet to that from supplements. Members noted that associations with haemorrhagic effects were reported following supplemental exposure to ginger, (Kruth *et al.*, 2003; Rubin *et al.*, 2019; Al Askar *et al.*, 2020) though these were inconclusive.

85. A study by ElMazoudy and Attia (2018) linked follicular failure to haemorrhagic effects in a study investigating the effect of dried ginger root on the oestrus cycle and implantation, in female mice. The authors concluded that ginger impairs the normal growth of the corpus luteum because of progesterone insufficiency during early pregnancy and that the results suggested that ginger can disrupt the oestrous cycle and blastocyst implantation without teratogenesis. They considered the highest NOAEL to be 500 mg/kg bw. The COT noted that this might be worth further investigation. However, it was also noted that other

factors could be contributing to the results observed and the study results were inconclusive.

### **In vitro studies**

86. Srivastava reported an effect of ginger extracts on *in vitro* platelet aggregation (1986). Ginger extracts in water, n-hexane, chloroform, and ethyl acetate were shown to inhibit platelet aggregation using arachidonic acid (AA), epinephrine, adenosine diphosphate (ADP), and collagen as agonists.

### **Animal studies**

87. The effect of an aqueous ginger extract on platelet thromboxane-B<sub>2</sub> (TXB<sub>2</sub>) and prostaglandin-E<sub>2</sub> (PGE<sub>2</sub>) production was studied by Thomson *et al.* (2002). Adult female Sprague-Dawley rats were administered an aqueous extract of raw ginger at either 50 mg/kg or 500 mg/kg daily, by either oral gavage or intraperitoneally (IP) for a period of 4 weeks. A dose of 50 mg/kg ginger administered orally, or IP did not result in any significant reduction in serum thromboxane-B<sub>2</sub> levels when compared to saline-treated control groups but doses at 500 mg/kg significantly reduced TXB<sub>2</sub> levels in serum.

88. A non-significant reduction in the level of TXB<sub>2</sub> was observed when ginger was injected IP. However, levels were not significantly different from the TXB<sub>2</sub> levels in control rats that had received saline. 50 mg/kg of ginger administered orally resulted in serum PGE<sub>2</sub> levels being significantly reduced however, 500 mg/kg was found to be more effective in reducing PGE<sub>2</sub> synthesis. PGE<sub>2</sub> levels were reported to be significantly lower than the saline control in rats given 500 mg/kg ginger extract both orally and IP.

89. The composition of ginger extracts also appears to vary according to whether ginger is fresh or dried. Suekawa *et al.*, demonstrated that (6)-shogaol, a principal component mainly found in dried ginger, inhibited carrageenan-induced swelling of rat hind paw, AA-induced platelet aggregation in rabbit and prostaglandin PGI<sub>2</sub> release in rat aorta, suggesting a potential inhibitory action on cyclooxygenases (COX) in both platelets and aorta tissue (1986, abstract only).

### **Human studies**

90. Srivastava (1989) investigated the effect of fresh ginger on blood platelet thromboxane synthesis in humans. In a study on 7 women aged between 25-65 years, where volunteers consumed ~5g of fresh ginger for 7 days. Ginger

was found to inhibit eicosanoid biosynthesis *in vivo*.

91. Krüth *et al.* reported the possible over-anticoagulation resulting from a possible ginger-phenprocoumon interaction (2003). A 76-year-old woman on long-term phenprocoumon therapy presented with epistaxis and an international normalized ratio (INR) of >10. Partial thromboplastin time (PTT) was also found to be prolonged (84.4 seconds; normal 35). For several weeks prior to the event, the woman had a regular ginger intake of dried ginger pieces and tea from ginger powder. Following treatment with vitamin K, the patient's INR and PTT returned to within therapeutic range.

92. Young *et al.* investigated the synergistic effect of ginger and nifedipine on anti-platelet aggregation in healthy volunteers aged 25-60 years old and hypertensive individuals aged 35-60 years old (2006). In a five-part study, the two groups comprising of 10 males and 10 females were administered 75 mg of acetylsalicylic acid (ASA), 1 g of ginger, 10 mg nifedipine, 1 g ginger and 10 mg nifedipine in combination and 1 g ginger and 75 mg ASA in combination daily for one week each following a washout period (7 days following ASA administration, 10 days thereafter).

93. Platelet aggregation in the presence of collagen ADP and epinephrine was 44.1%, 44.5% and 42.1% in normal subjects and 64.2%, 67.7% and 62.9% in hypertensive patients, respectively. Platelet aggregation induced by collagen, ADP or epinephrine was found to be higher in hypertensive patients than normal patients. Following administration of ginger alone, platelet aggregation was measured as 35.2%, 37.8%, 35.9% with collagen, ADP and epinephrine respectively. When administered ginger and nifedipine in combination, the percentage inhibition of platelet aggregation induced by collagen, ADP and epinephrine were 79.8%, 75.2%, 69.3% respectively.

94. Rubin *et al.* (2019) reported the possible effect of ginger supplementation on the (INR) in a woman taking warfarin. The 70-year-old female, who had been taking clonazepam 1 mg, metoprolol succinate 25 mg, paroxetine 10 mg, phenytoin 30 mg, rosuvastatin 20 mg, warfarin 7.5 mg daily, and warfarin 10 mg once day per week, presented with an INR of 8, an increase from 2.7 one month prior after taking a 48 mg ginger supplement daily. A week following cessation of the ginger supplement, the INR declined to 2.6.

95. Al Askar *et al.* (2020) investigated the effect of ginger on platelet aggregation using agonists adenosine diphosphonate, arachidonic acid, collagen, ristocetin and epinephrine. Forty healthy male and female participants were

randomized (1:1) to consume ginger tea at an amount of 4 g in 150 ml of boiling water once daily vs. 4 g twice daily for five consecutive days. 4 g of ginger powder administered daily resulted in reduced platelet aggregation in subjects using epinephrine only. Platelet aggregation inhibition was also found to be higher in women using arachidonic acid.

96. Ginger, in powder form (5 g per day), was demonstrated to significantly (P0.001) decrease ADP- and epinephrine-induced platelet aggregation in healthy male subjects who were fed 100 g of butter daily for seven days (Verma *et al.*, 1993). Conversely, Bordia *et al.*, found that 4 g powdered ginger administered daily over the course of 1.5 and 3 months had no effect on ADP and epinephrine-induced platelet aggregation in individuals with coronary artery disease (CAD). However, a single 10g dose of powdered ginger, administered to CAD patients resulted a significant decrease in induced platelet aggregation (1997).

97. Conversely, Lumb found that a dose of 2g of ginger in powder form daily produced no significant differences in platelet aggregation/function than the placebo and concluded that these effects may be dose dependant or attributed to fresh ginger (1994). Whilst investigating the effects of ginger on the pharmacokinetics or pharmacodynamics of warfarin and the effect of ginger on clotting status in adult males, Jiang *et al.*, also found that the pharmacokinetics nor pharmacodynamics of warfarin were not affected in healthy males who were treated with a single 25 mg dose of warfarin, following 7 days of pretreatment with ginger tablets (3 tablets, 3 times per day, each capsule containing extract equivalent to 0.4 g of ginger rhizome powder) (2005). Furthermore, ginger had no effect on international normalized ratio (INR) or *ex vivo* platelet aggregation in response to arachidonic acid.

## **Effects on blood pressure**

### **In vivo**

98. Ghayur and Gilani (2005) reported that a crude extract of ginger induced a dose-dependent (0.3–3 mg/kg) fall in arterial blood pressure of anesthetized Sprague-Dawley rats (2005). In Guinea pig paired atria, the crude extract exhibited cardio-depressant activity on the rate and force of spontaneous contractions. In rabbit thoracic aorta preparation, the crude extract relaxed the phenylephrine induced vascular contraction at a dose 10 times higher than that required against K<sup>+</sup>-induced contraction (80 mM).

99.  $\text{Ca}^{2+}$  channel-blocking (CCB) activity was confirmed when the crude extract shifted the  $\text{Ca}^{2+}$  dose-response curves to the right similar to the effect of verapamil. It also inhibited the phenylephrine (1 mM) control peaks in normal- $\text{Ca}^{2+}$  and  $\text{Ca}^{2+}$ -free solution, indicating that it acts at both the membrane-bound and the intracellular  $\text{Ca}^{2+}$  channels. When tested in endothelium-intact rat aorta, it again relaxed the  $\text{K}^{+}$ -induced contraction at a dose 14 times less than that required for relaxing the PE-induced contraction. The vasodilator effect of the crude extract was endothelium-independent because it was not blocked by N $\omega$ -nitro-L-arginine methyl ester hydrochloride (L-NAME) (0.1 mM) or atropine (1 mM) and also was reproduced in the endothelium-denuded preparations at the same dose range. These data indicate that the blood pressure-lowering effect of ginger is mediated through blockade of voltage-dependent calcium channels.

## Effect on Prostaglandins

### In vitro

100. Ginger extracts, along with many gingerols and shogaols have been shown to suppress prostaglandin synthesis *in vitro*, through inhibition of cyclooxygenase (Jolad et al. 2005; Pan et al. 2008; Dugasani et al. 2010).

101. Lantz *et al.* investigated the anti-inflammatory effect of ginger extracts and the principal components thereof (6-, 8- 10-gingerols and 6-, 8-, 10-shogaols) in an *in vitro* model. U937 cells, differentiated and exposed to lipopolysaccharide (LPS) from *Escherichia coli* (1 mg/ml). Extracts containing predominantly gingerols were found not to be cytotoxic, while shogaols were found to be cytotoxic at concentrations above 20  $\mu\text{g}/\text{ml}$  (2007).

102. Jolad *et al.* also demonstrated the inhibitory effect of gingerols on LPS-induced PGE<sub>2</sub> production in HL-60 cells stimulated with 1  $\mu\text{g}/\text{ml}$  of LPS (2004). None of the compounds tested were shown to be cytotoxic.

## Contaminants

103. Differences in cultivation conditions and extraction methods could lead to possible sources of contamination from toxins, microbes, pesticides, heavy metals and residual solvents. Studies investigating contamination in ginger are limited, however of the few available, the main sources of contamination reported are heavy metals (Wagesho & Chandravanshi, 2015; Goroya *et al.*, 2019, Kilic & Soylak, 2019; Xu *et al.*, 2020) and mycotoxins (Ałtyn and Twarużek, 2020; Wen *et*

*al., 2014; Omotayo et al., 2019; Lippolis et al., 2017).*

104. Ginger can be exposed to mycotoxin contamination during harvesting, storage and handling. Whilst information on mycotoxin contamination in ginger is limited, ginger has been demonstrated to be particularly exposed to aflatoxins and ochratoxin A (OTA). This is reflected in GB legislation where maximum levels for spices including ginger are established in [Retained EU Law 1881/2006](#). Maximum levels of 5 µg/kg for aflatoxin B1 (AFB1), 10 µg/kg for all aflatoxins (sum of AFB1, AFB2, AFG1, and AFG2) and 15 µg/kg for OTA, for ginger and its products.

105. The Committee discussed the potential presence of contaminants in ginger and noting that the ginger products used in the studies reported were sourced locally in markets or herbalists (Wagesho & Chandravanshi, 2015; Goroya *et al.*, 2019), Members queried whether there were any specific data on contaminants in ginger supplements available in the UK.

106. The Committee noted it was unknown how much ginger and particularly, highly concentrated juice extracts, would contribute to overall contaminant exposure in the UK.