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1 **ABSTRACT**

2 **Objectives:**

3 Exposure to sex hormones is important in the pathogenesis of breast cancer (BC) and inability to  
4 tolerate such exposure may be reflected in increased asymmetrical growth of the breasts. This study  
5 aims to characterise, for the first time, asymmetry in breast volume (BV) and radio-dense volume  
6 (DV) in a large ethnically-diverse population.

7 **Methods:**

8 Automated measurements from digital raw mammographic images of 54,591 cancer-free  
9 participants (aged 47-73) in a UK breast screening programme were used to calculate absolute (cm<sup>3</sup>)  
10 and relative asymmetry in BV and DV. Logistic regression models were fitted to assess asymmetry  
11 associations with age and ethnicity.

12 **Results:**

13 BV and DV absolute asymmetry were positively correlated with the corresponding volumetric  
14 dimension (BV or DV). BV absolute asymmetry increased, whilst DV absolute asymmetry decreased,  
15 with increasing age (P-for-linear-trend<0.001 for both). Relative to Whites, Blacks had statistically  
16 significantly higher, and Chinese lower, BV and DV absolute asymmetries. However, after adjustment  
17 for the corresponding underlying volumetric dimension the age and ethnic differences were greatly  
18 attenuated. Median relative (fluctuating) BV and DV asymmetry were 2.34% and 3.28% respectively.

19 **Conclusions:**

20 After adjusting for the relevant volumetric dimension (BV or DV), age and ethnic differences in  
21 absolute breast asymmetry were largely resolved.

22 **Advances in knowledge:**

23 Previous small studies have reported breast asymmetry – BC associations. Automated  
24 measurements of asymmetry allow the conduct of large-scale studies to further investigate these  
25 associations.

26

27

## 28 INTRODUCTION

29 Exposure to endogenous and exogenous sex hormones are recognized to be important in breast  
30 development and in the pathogenesis of breast cancer [1-5], with the effect of many reproductive  
31 factors on breast cancer risk, e.g. early age at menarche and late age at menopause, being mediated  
32 by circulating levels of these hormones [6]. There is also some evidence that pre-natal exposure to  
33 high levels of sex hormones may increase the risk of breast cancer. Breast cancer risk is elevated in  
34 women who were exposed in utero to diethylstilboestrol (DES) given to their mothers to prevent  
35 pregnancy complications [7] and some studies have reported positive associations between breast  
36 cancer risk and birth size, pre-eclampsia and multiple births, all possible markers of raised, in-utero,  
37 exposure to oestrogens [8]. It is also thought that an individual's ability to tolerate exposure to  
38 oestrogens, particularly during periods of growth, may be reflected in a higher degree of  
39 homeostasis and thus bilateral symmetrical development of paired organs such as the breasts [9].  
40 Increased 'fluctuating asymmetry', i.e. increased anthropometrical asymmetry in paired features, is  
41 a common response to increased stress during development [10] and is related to both fecundity  
42 and general health [11-14]. For example, studies of dermatoglyphics have shown that increased  
43 asymmetry in hand patterns is associated with increased risk of several diseases including breast  
44 cancer [15]. Also, women with high 2<sup>nd</sup> digit to 4<sup>th</sup> digit ratio (2D:4D) (thought to be associated with  
45 lower exposure or sensitivity to prenatal testosterone and/or higher levels in utero oestrogen levels)  
46 had increased risk of breast cancer [16] and they presented with breast cancer at a younger age [17,  
47 18]. An association between left-handedness and increased risk of breast cancer has also been  
48 reported [19, 20]. Manning et al showed that increased breast FA was correlated not only with age,  
49 height and parenchymal type but also with reproductive factors such as parity, age at first birth and  
50 age at menopause [9].

51 Only a few small-sized studies, mainly among Caucasians, have so far examined the association  
52 between breast size asymmetry and breast cancer risk. Their findings are consistent with asymmetry  
53 being associated with the presence of a breast cancer [21-24] as well as with a higher risk of having a  
54 breast cancer diagnosed in the short- and medium-term (mean interval between mammography  
55 and diagnosis 6.44 years) [25]. Mammographic density captures the amount of radio-dense tissue in  
56 the breast, and there is also some evidence that asymmetry in density might be associated with  
57 higher short-term likelihood of being diagnosed with breast cancer [26-28]. It has also been  
58 suggested that a slightly larger left breast, with a higher volume of radio-dense tissue, may account  
59 for the slightly higher frequency of cancers in the left than the right breast although the mechanisms  
60 for this are poorly understood [29-31]. Overall, the findings from these studies suggest that  
61 asymmetry in breast size and density may reflect underlying biological mechanisms linked to the  
62 pathogenesis of breast cancer or may be early consequences of the presence of a tumour. Hence,  
63 asymmetry measurements have the potential to be used as risk predictors or diagnostic markers. To  
64 our knowledge there is, as yet, no large-scale study of the prevalence of breast volume asymmetry  
65 and breast density asymmetry from large population-based studies.

66 The recent introduction of full-field digital mammography (FFDM) has led to the development of  
67 automated algorithms which allow volumetric assessments of both breast size and mammographic  
68 density from 2-dimensional digital mammographic images. Such automated methods make it  
69 feasible to conduct large-scale studies based on objective measurements of bilateral asymmetry in  
70 breast size and mammographic density. This study aims to quantify bilateral asymmetry in breast

71 size and mammographic density volume in a very large, and ethnically-diverse sample of over 54,000  
72 women who participated in a population-based breast screening programme in England. The  
73 findings will provide the first population-based data on the distribution of breast asymmetry, and  
74 potential age and ethnic variations.

## 75 **METHODS**

### 76 **Study participants**

77 The study participants were women resident in one of five London boroughs – Wandsworth,  
78 Merton, Croydon, Sutton, Richmond and Kingston – who underwent routine 3-yearly screening  
79 mammography as part of the England and Wales National Health Service Breast Screening  
80 Programme (NHSBSP) at the South West London Breast Screening Service (SWLBSS) based in the St  
81 George's University Hospitals National Health Service (NHS) Foundation Trust. The NHSBSP is an  
82 organised population-based mammographic screening programme, with a call-recall system, which  
83 targets women aged 50-70 years and has a coverage of ~75% [32]. Also included were a small  
84 number of younger women (aged 29-45) who had been identified as having a higher risk of breast  
85 cancer and therefore were invited for screening on an annual basis [33], plus any women over 73  
86 years who had optionally contacted the service for a self-referred screening appointment. All  
87 women were asymptomatic at the time of screening. Participants were screened during the period  
88 01/03/2013 to 18/08/2016. Data on ethnicity were collected as part of the standard screening  
89 protocol via a self-completed screening questionnaire. Ethnicity was categorised according to the  
90 Census classification and summarised as, "Asian" (Indian, Pakistani or Bangladeshi or other), "Black-  
91 African", "Black-British or Caribbean or other", "Chinese", "Mixed" (White and Black, White and  
92 Asian or any other mixed), "White" (British or Irish or other) and "Other" [34]. Data for other known  
93 breast cancer risk factors (e.g. parity, duration of breast feeding, age at menarche, body mass index  
94 (BMI), family-history of breast cancer) are not collected in a systematic way across the NHSBSP  
95 screening programme and thus were unavailable.

96  
97 Each woman underwent the NHSBSP standard 2-view (cranio-caudal (CC) and medio-lateral-oblique  
98 views (MLO)) mammography of each breast [35], with the set of four digital raw images being stored  
99 on the SWLBSS Picture Archiving and Communication system (PACS). The images were double read  
100 with arbitration by consensus. When women had multiple screening episodes during the study  
101 period, only images from the earliest screen episode were included in the analysis. Raw digital  
102 mammographic images were processed via the automated algorithm Volpara® Density™ version  
103 1.5.11 (Volpara), (Matakina Technology Limited, Wellington, New Zealand) [36]; this algorithm  
104 provided fully-automated estimates (in cm<sup>3</sup>) of the volume of the breast (BV) and the volume of the  
105 radio-dense tissue (DV) separately for each of the four (left (L) and right (R) breasts / CC and MLO  
106 views) images. The screening programme does not use mammographic density as a diagnostic aid,  
107 and participants are not informed on whether they have dense breasts.

108 In all, 66,176 women were screened during the study period. Women were excluded from this  
109 analysis if cancer was detected by the current screen (N=530); if they had a previous history of  
110 breast cancer (N=438); if their screen images were classified as "technical recall", i.e. were  
111 considered by the reader not to be of high enough quality for diagnosis (N=26); if they had breast  
112 implants; if their standard set of four images (i.e. L/R CC and MLO images) was incomplete

113 (N=9,823); and if at least one of the two CC images was rejected by Volpara based on its internal  
114 consistency checks (N=7,338). Exclusions were not mutually exclusive, leaving a total of 54,591  
115 women who were eligible for inclusion in the analysis.

## 116 **Ethical approval**

117 This retrospective study was carried out on fully anonymous, routinely collected data only, held in  
118 accordance with the NHS Cancer Screening Programmes Confidentiality and Disclosure Policy 2011.  
119 The NHSBSP has section 251 support under the NHS Act 2006. The study was approved by all  
120 relevant ethics committees (Research Ethics Committees from St George's University Hospitals NHS  
121 Foundation Trust, and the London School of Hygiene and Tropical Medicine).

## 122 **Statistical Methods**

123 For each participant the volume of each breast (BV), and the volume of radio-dense tissue (DV), was  
124 calculated as the average of the readings obtained from the same side CC and MLO images (i.e. CC  
125 and MLO views were used to obtain an overall average). Both absolute and relative measures of left-  
126 right asymmetry were calculated: *absolute* asymmetry (in  $\text{cm}^3$ ), i.e. the unsigned difference between  
127 left BV (or DV) and right BV (or DV), and *relative* asymmetry as  $(|L-R|)/(L+R)/2$  expressed as a  
128 percentage. Absolute and relative asymmetry were estimated from the CC images only because this  
129 view is likely to capture the whole of the breast whilst being less affected than the MLO view by the  
130 inclusion of variable amounts of retro-glandular fat tissue near the chest wall [36]. (For comparison  
131 the equivalent asymmetry measures were also calculated using the MLO views only).

132 The distributions of absolute and relative asymmetry values were plotted. Natural-log  
133 transformations were applied to normalise the distributions of absolute and relative BV and DV  
134 asymmetry and quintiles were used to categorise BV and DV into five equally sized categories.

135 To examine whether age-related variations in breast volume and breast asymmetry differ across the  
136 various ethnic groups, medians, 25<sup>th</sup> and 75<sup>th</sup> centiles of the distributions of untransformed BV, DV  
137 and absolute asymmetry measures were also calculated and plotted separately by 5-year age  
138 categories and ethnicity. These were also calculated for each single year of age and plotted after  
139 smoothing using a Lowess function (values based on fewer than 20 observations were omitted from  
140 the plots). Scatter plots and Spearman correlation coefficients were used to examine the  
141 correlations between asymmetry measures and the corresponding volumetric dimension. In order to  
142 assess whether allometry is a feature of this relationship (as identified by Manning et al [9]) we  
143 regressed log of asymmetry on log of the corresponding volumetric measure.

144 Linear regression models were used to examine the strength of the associations between each  
145 exposure variable – age and ethnicity – and the outcome variables, BV or DV absolute asymmetry,  
146 controlling for their respective average volume (BV or DV). Because of the log-transformation,  
147 regression coefficients represent the relative change (RC) in absolute asymmetry per one unit  
148 change in the exposure category. In all the analyses, we considered statistical significance (2-sided)  
149 at  $p\text{-value} < 0.05$ . All analyses were conducted in Stata (IC 14) [37].

150 **RESULTS**

151 **Study participants**

152 The characteristics of the 54,591 participants are shown in Table 1. The majority (~87%) of women  
153 were within the ages of 50 to 70 years, the age-group targeted by the NHSBSP. Among the 85% of  
154 the participants who reported their ethnicity, ~76% were White but there were also high numbers of  
155 women of Black and Asian ethnicity.

**Table 1 Characteristics of the study participants**

	No.	Percent	BV (cm <sup>3</sup> ) <sup>a</sup>	DV (cm <sup>3</sup> ) <sup>a</sup>	Median (25 <sup>th</sup> and 75 <sup>th</sup> centiles)			
					BV Absolute CC Asymmetry <sup>b</sup> (cm <sup>3</sup> )	DV Absolute CC Asymmetry <sup>b</sup> (cm <sup>3</sup> )	BV Relative CC Asymmetry (%) <sup>c</sup>	DV Relative CC Asymmetry (%) <sup>c</sup>
<b>Age at screening (yrs)</b>								
<45-	234	0.4	563 (353, 950)	63.8 (47.4, 94.7)	56.7 (24.7, 105.6)	7.79 (3.86, 16.18)	2.87 (1.48, 4.56)	3.26 (1.87, 5.74)
45-49-	3,297	6.0	727 (450, 1135)	62.0 (45.4, 85.3)	57.5 (24.7, 112.8)	7.46 (3.39, 14.46)	2.42 (1.11, 4.06)	3.40 (1.62, 5.83)
50-54	15,40	28.2	762 (485, 1138)	54.3 (40.6, 75.5)	59.2 (25.8, 115.4)	6.45 (2.79, 12.69)	2.36 (1.12, 4.09)	3.33 (1.54, 5.87)
55-59	12,40	22.7	770 (498, 1148)	48.5 (36.6, 64.6)	61.8 (26.8, 120.4)	5.59 (2.45, 10.86)	2.43 (1.15, 4.21)	3.26 (1.53, 5.82)
60-64	10,44	19.1	767 (515, 1109)	46.1 (35.1, 61.0)	60.1 (26.9, 117.1)	5.21 (2.26, 10.24)	2.41 (1.14, 4.13)	3.18 (1.47, 5.65)
65-69	9,483	17.4	751 (506, 1063)	44.0 (33.9, 57.7)	62.5 (27.5, 120.9)	5.04 (2.19, 10.04)	2.50 (1.17, 4.33)	3.23 (1.47, 5.78)
70+	3,297	6.0	723 (499, 1014)	42.9 (33.5, 56.1)	63.6 (28.3, 118.9)	5.25 (2.27, 10.10)	2.66 (1.24, 4.52)	3.28 (1.52, 5.79)
Missing	27	0.1						
<b>Ethnic group</b>								
White - British, Irish, Other	35,44	64.9	747 (485, 1098)	47.9 (36.1, 64.9)	59.3 (25.8, 115.5)	5.60 (2.44, 11.13)	2.42 (1.13, 4.18)	3.30 (1.53, 5.84)
Asian <sup>d</sup>	4,829	8.9	718 (508, 1005)	44.8 (34.8, 59.6)	59.4 (27.2, 111.8)	5.02 (2.10, 9.91)	2.43 (1.19, 4.24)	3.17 (1.40, 5.52)
Black – British, Caribbean	2,705	5.0	956 (610, 1381)	58.3 (44.6, 77.8)	71.6 (31.1, 136.4)	6.59 (3.02, 12.17)	2.26 (1.04, 4.02)	3.17 (1.50, 5.49)
Black – African	1,999	3.7	960 (672, 1347)	56.0 (42.1, 74.0)	81.1 (35.7, 155.5)	6.39 (2.95, 12.65)	2.50 (1.20, 4.14)	3.23 (1.50, 5.64)
Mixed <sup>e</sup>	1,029	1.9	800 (535, 1176)	53.0 (39.4, 71.5)	64.5 (28.4, 124.5)	6.12 (2.71, 11.66)	2.36 (1.13, 4.21)	3.28 (1.50, 5.54)
Chinese	654	1.2	394 (258, 552)	41.0 (29.6, 60.7)	35.1 (16.2, 67.7)	5.03 (2.28, 9.67)	2.71 (1.38, 4.68)	3.38 (1.60, 6.52)
Missing or not reported	7,932	14.5	751 (499, 1121)	51.2 (38.5, 70.8)	61.6 (27.5, 119.5)	6.20 (2.71, 12.05)	2.48 (1.19, 4.21)	3.35 (1.56, 5.91)
<b>All women</b>	<b>54,59</b>		<b>757 (496, 1112)</b>	<b>48.9 (36.8, 66.5)</b>	<b>60.6 (26.6, 117.8)</b>	<b>5.71 (2.49, 11.27)</b>	<b>2.43 (1.15, 4.19)</b>	<b>3.28 (1.52, 5.79)</b>

Footnotes:

<sup>a</sup> Calculated from the average BV (or DV) value from the 4 images: left CC image, right CC image, left MLO image, right MLO image.

<sup>b</sup> Calculated as the absolute difference between the BV (or DV) value from the left CC image and the BV (or DV) value from the right CC image.

<sup>c</sup> Relative Asymmetry estimated as  $(|L-R|)/(L+R)/2*100$ , where L and R are volumes from the left and right breasts estimates from the CC views.

<sup>d</sup> Asian includes: British Indian, Pakistani, Bangladeshi, Other Asian excluding Chinese

<sup>e</sup> Mixed includes: White and Black, White and Asian or any other mixed



162 **Breast volume, dense volume and absolute asymmetry by age and ethnicity**

163 The median (25<sup>th</sup>, 75<sup>th</sup> centiles) BV and DV values for the whole study sample were 757 (496, 1112)  
164 cm<sup>3</sup> and 48.9 (36.8, 66.5) cm<sup>3</sup>, respectively (Table 1). There was, however, evidence of bilateral  
165 asymmetry in BV and DV, with a median (25<sup>th</sup>, 75<sup>th</sup> centiles) absolute difference in BV and DV  
166 between the two breasts of 60.6 (26.6, 117.8) cm<sup>3</sup> and 5.71 (2.49, 11.27) cm<sup>3</sup>, respectively, with the  
167 wide IQR indicating considerable between-woman variation in bilateral asymmetry (Table 1). This  
168 difference was seen in every age and ethnic group, albeit with some variations with the smallest  
169 median absolute differences seen among Chinese women.

170 The distributions of BV and DV absolute asymmetry estimates were right skewed and, hence, a log-  
171 normal transformation was used to normalise them (Figure 1). The transformed BV and DV  
172 asymmetry distributions approximated a normal distribution although both were leptokurtic  
173 (kurtosis coefficient: 5.60 and 4.76, respectively) and slightly skewed (skewness coefficient: -1.12  
174 and -0.96, respectively).

175 Further analyses by age-group show that, on average, BV increased slightly with increasing age up to  
176 ages 55-59, declining thereafter (Figure 2). Ethnic variations in BV were much more marked than  
177 those observed with age (Figure 3), with BV being, on average, highest among Black Caribbean  
178 (median: 956 cm<sup>3</sup>) and Black African (960 cm<sup>3</sup>) women and lowest among Chinese women (394 cm<sup>3</sup>)  
179 but with wide between-woman variability being present within each ethnic group. Absolute BV  
180 asymmetry showed similar age and ethnicity patterns to those observed for BV (Figures 2 and 3).

181 In contrast to BV, DV decreased, on average, with increasing age-group from <45 to 70+ years but,  
182 similarly to BV, DV was highest among Black Caribbean (median: 58.3 cm<sup>3</sup>) and Black African women  
183 (56.0 cm<sup>3</sup>) and lowest among Chinese women (41.0 cm<sup>3</sup>). Absolute DV asymmetry followed a similar  
184 pattern to DV, i.e. lower values across successive age-groups, and higher among Black African and  
185 Black Caribbean women (Figures 2 and 3).

186 The observed absolute asymmetry in BV and DV reflected that fact that, on average, women had a  
187 larger left breast with a larger amount of radio-dense tissue. The only exception was that DV was  
188 higher in the right breast among Chinese women.

189 Figure 4, which depicts median single-year-of-age volumetric and asymmetry values by ethnicity,  
190 shows that age-related changes in BV varied across the different ethnic groups. Among Asian, Black  
191 African and White women, BV increased progressively up to age ~60 years but declined thereafter  
192 whilst among Black Caribbean women, BV continued to increase up to age 70 years. In contrast, DV  
193 decreased with age in all ethnic groups. There was, however, a marked levelling out after age ~55.  
194 BV and DV absolute asymmetry follow the same general pattern as their corresponding underlying  
195 volumetric dimension.

196 **Relative asymmetry by age and ethnicity**

197 The magnitude of relative BV asymmetry was similar across all age groups (median overall relative  
198 BV asymmetry for all study participants: 2.43% (25<sup>th</sup>, 75<sup>th</sup> centiles: (1.15%, 4.19%); Table 1) except  
199 that it was slightly higher in the youngest age band (median 2.87% (1.48%, 4.56%)). The magnitude  
200 of relative BV asymmetry was also similar irrespective of the ethnicity of the participants although  
201 slightly higher in the Chinese ethnic group (2.71% (1.38%, 4.68%)).

202 The magnitude of relative DV asymmetry was similar across all age groups and ethnicities (median  
203 overall relative DV asymmetry for all study participants: 3.28% (1.52%, 5.79%)). Overall age and  
204 ethnic variations in relative BV and DV asymmetry were much less marked than those observed for  
205 absolute BV asymmetry and absolute DV asymmetry (Figures 2 and 3).

#### 206 **Correlations between absolute asymmetry and volumetric measures**

207 BV and DV absolute asymmetry were moderately positively associated with their corresponding  
208 underlying volumetric measure (Spearman correlation coefficient (r): 0.45 and 0.43, respectively;  
209  $P < 0.0001$  for both). Regressing log BV asymmetry on log BV revealed negative allometry (coefficient:  
210 0.84; 95% CI 0.83, 0.85)) whilst regressing log DV on log DV revealed slight positive allometry (1.09;  
211 1.07, 1.12). There were no statistically significant differences in the magnitude of these allometry  
212 coefficients across the different ethnic groups (data not shown).

#### 213 **Associations between absolute asymmetry and age and ethnicity**

214 The fitted linear regression models showed that BV absolute asymmetry increased with increasing  
215 age (in 5-year categories, P for trend (Pt) $< 0.001$ ; Table 2), and that this trend persisted after  
216 adjustment for BV (Pt $< 0.001$ ). In contrast, DV absolute asymmetry decreased with increasing age  
217 (Pt $< 0.001$ ), but this trend was attenuated upon adjustment for DV (Pt=0.14; Table 2). Further  
218 adjustment for ethnicity affected little the magnitude of the BV or DV absolute asymmetry  
219 associations with age (Table 2).

220 When considering ethnicity on its own, relative to White women (reference group) those of Black  
221 Caribbean, Black African and Mixed ethnicity had statistically significantly higher, whilst those of  
222 Chinese ethnicity had statistically significant lower, BV absolute asymmetry (Table 2). However, upon  
223 adjustment for BV the magnitude of these ethnic differentials was markedly reduced, remaining  
224 statistically significant only in Black African women (RC 1.13; 95% CI 1.07, 1.19), while there was  
225 borderline evidence of higher BV absolute asymmetry for Asian women (1.04; 1.00, 1.07; Table 2).  
226 Similarly, and still relative to White women, DV absolute asymmetry was found to be significantly  
227 higher among Black Caribbean and Black African women and significantly lower among Asian and  
228 Chinese women in unadjusted analyses. However, these differences remained significant after,  
229 adjustment for DV, only for Asian women (0.94; 0.91, 0.98; Table 2). There was no evidence of  
230 interaction between age and ethnicity in their effects on BV or DV absolute symmetry ( $p = 0.69$  and  
231  $p = 0.53$ , respectively).

## 232 **DISCUSSION**

### 233 **Main findings**

234 This study of >54,000 women clarifies the associations between absolute breast asymmetry and  
235 breast volume, with the findings being broadly consistent with those from a smaller study (n=500  
236 younger women) by Manning et al. which showed that simple linear regression of BV absolute  
237 asymmetry (log transformed) on BV gives a significant positive association (our study  $r^2 = 0.15$ ,  
238  $p < 0.001$ ; Manning  $r^2 = 0.13$ ,  $p < 0.001$ ) [9]. We also found that absolute DV asymmetry is strongly  
239 positively associated with DV. Thus, the larger the volume of the breast (or the volume of the radio-  
240 dense tissue) the higher the magnitude of BV (or DV) absolute asymmetry. This explained, at least in  
241 part, the higher levels of BV and DV asymmetry observed in women of Black ancestry as they also

242 had, on average, higher BV and DV. After adjusting for the relevant breast volumetric measure (i.e.  
243 BV for BV asymmetry, DV for DV asymmetry), the ethnic differences in absolute breast asymmetry  
244 observed in the unadjusted analysis were attenuated, indicating that they were largely driven by  
245 ethnic differences in breast and dense tissue volumes.

246 Similar to the findings of Manning et al. [9], our findings showed that the BV absolute asymmetry/BV  
247 relationship was negatively allometric across all main ethnic groups, indicating that women with  
248 large breasts had a smaller fluctuating asymmetry than expected for their volume. There was,  
249 however, evidence that the DV absolute asymmetry/DV relationship was positively allometric.

250 Like Manning et al. we found, using simple linear regression, that BV asymmetry is only weakly  
251 positively associated with age (our study  $r^2=0.004$ ,  $p<0.001$ , Manning  $r^2=0.019$ ,  $p=0.02$ ) [9]. The  
252 differences in the strength of the association might be explained by the fact that the women in our  
253 study were considerably older than those in the study by Manning et al.[9] (mean ages 58.57 and  
254 39.85 respectively). We found that DV absolute asymmetry is weakly but negatively associated with  
255 age, with these associations being attenuated upon adjustment for DV, indicating that these  
256 associations are largely driven by decreasing DV with age.

257 Two earlier studies, one in the USA ( $n=980$ ) [38] and the other in Switzerland ( $n=87$ ) [39], focused on  
258 the left:right ratio (L:R) in BV. Although such L:R ratio cannot be regarded as a measure of relative  
259 asymmetry, it is nevertheless worth noting that their findings are consistent with our finding that, on  
260 average, the left BV exceed the right BV by ~4% across the whole breast screening population  
261 irrespective of ethnicity and age. There was, however, marked between-woman variability in breast  
262 asymmetry among cancer-free, screened women.

263 Literature on the prevalence of DV asymmetry is limited. Consistent with our findings Lee et al., in a  
264 study of 860 South Korean women, found that the L:R ratio in DV was less than 1 indicating a greater  
265 volume of radio-dense tissue in the right breast, thus challenging the view that the laterality of DV  
266 ratio is similar across all ethnic groups. Chen et al. [40] on a small sample of 24 Taiwanese women  
267 also found that DV, as measured by magnetic resonance imaging (MRI), was higher in the right than  
268 in the left breast.

## 269 **Strengths and limitations**

270 Strengths of this study include its population-based design, the very large sample size relative to  
271 previous studies, and the wide ethnic mix. As the images for both breasts were collected at the same  
272 point in time, and under similar technical conditions, within-woman L:R breast comparisons are  
273 unlikely to have been biased by anthropometric, reproductive and lifestyle characteristics (e.g. BMI,  
274 menopausal status) or by differences in image acquisition (e.g. differences in mammographic  
275 equipment) as these would have affected both breasts similarly. This does not exclude, however, the  
276 possibility that the findings may have been affected by within-woman differences in the way the left  
277 and right breasts were examined (e.g. differences in a woman's positioning during mammography).  
278 The study relied on an automated method to estimate the volumes of the left and right breasts and  
279 the amounts of their radio-dense tissues, and thus such objective measurements were not  
280 influenced by subject or observer biases. Although the volumetric estimates were derived from 2-  
281 dimensional images and, hence, may have been affected by errors, these would have affected both  
282 breasts similarly.

283 The study included mostly women of screening age and reflected a mix of ethnic groups living in  
284 England. The proportion (15%) of women for whom ethnicity data were missing was relatively low  
285 and typical for NHSBSP screening services where collection of self-reported ethnicity data is  
286 undertaken [42]. Women with a previous history of breast cancer, or who were diagnosed with  
287 cancer at the time of screening, as well as those with breast implants, were excluded from the  
288 study; however, women with other conditions that might have affected their breast size (e.g. surgery  
289 for non-malignant conditions) could not be excluded as information on these conditions is not  
290 routinely collected by the NHSBSP.

291 A limitation of this study was the lack of data on potential confounders or mediators (e.g. BMI,  
292 reproductive history) of the age/ethnicity associations with BV and DV asymmetry. Menstrual cyclic  
293 variations in breast width asymmetry (measured from CC mammograms) were reported by Manning  
294 et al [43], based on mammograms from 280 premenopausal women, with lowest breast asymmetry  
295 occurring around the middle of the cycle (which Scutt & Manning later attributed to ovulation [44]).  
296 Although the present study was unable to consider cyclical changes in asymmetry as information on  
297 the day of menstrual cycle when the mammogram was taken is not routinely collected by the  
298 NHSBSP, the large majority of women screened by the NHSBSP are of postmenopausal age.  
299 Nevertheless, future studies of pre-menopausal women should examine cyclic variations in  
300 asymmetry and, in particular, whether such variations should be taken into account when assessing  
301 asymmetry – breast cancer risk associations.

302 The study was conducted using one specific algorithm for estimating volumetric breast size and  
303 volumetric density. There is no published data specifically on the reliability of asymmetry measures  
304 derived from the Volpara volumetric measurements, but the latter have been found to be reliable  
305 and repeatable [45-47]. Nevertheless, it would be worthwhile to assess breast asymmetry using  
306 other automated methods. Our estimates of BV and DV asymmetry were derived from the CC views  
307 of the left and right breasts; however, MLO views produced similar breast asymmetry estimates (e.g.  
308 median (IQR) for BV and DV absolute asymmetry for all participants was 60.6 (26.6, 117.8) cm<sup>3</sup> and  
309 5.71 (2.5, 11.3) cm<sup>3</sup>, respectively, if derived from the CC views and 65.1 (28.7, 127.0) cm<sup>3</sup> and 7.2  
310 (3.2, 14.1) cm<sup>3</sup>, respectively, if derived from the MLO views). Similar associations of these measures  
311 with age and ethnicity were also found (data not shown).

## 312 **Implications**

313 So far, only a few small, studies have examined the relation of breast asymmetry measures with  
314 breast cancer. Scutt et al. used area-based mammographic breast size (BV) asymmetry  
315 measurements from ~250 breast cancer cases and ~250 matched controls, while adjusting for known  
316 risk factors and absolute breast size, to show that absolute BV asymmetry at baseline screen was  
317 associated, with cancer diagnosis at the baseline screen [21] and also medium-term risk [22]. In a  
318 preliminary study, Eltonsy et al. examined data from 280 breast cancer cases and 82 controls and  
319 found that the mean absolute BV asymmetry, adjusting for BV, was significantly higher in cancer  
320 patients [19]. Kayar et al. used non-mammographic breast measurements (from Grossman-  
321 Rounder Discs) on 251 breast cancer cases and 466 controls from a Turkish outpatient clinic, to  
322 propose a 'pathological breast asymmetry ratio', suggesting that a L:R BV ratio of >±20% was  
323 associated with an increased risk of breast cancer being diagnosed within one year of the  
324 examination [20].

325 Zheng et al. investigated the relationship between mammographic density percentage (%MD)  
326 asymmetry and breast cancer using a bespoke algorithm on mammograms from 230 women with  
327 interval cancers (cancers diagnosed between screens) and 230 controls and suggested that as  
328 percent mammographic density (i.e. volume of radio-dense tissue measured as percent of the total  
329 breast volume) asymmetry increases there was an increased risk of cancer at both current screen  
330 and in the medium term (1-3 years). These models adjusted for subjective breast density category  
331 (BIRADS), but not for absolute breast density [23, 24].

332 The limited available literature suggests that BV and DV asymmetry may have potential value as  
333 markers of either the presence of a cancer (diagnostic marker) or the risk of developing cancer in the  
334 future (risk predictor). Proper examination of the potential value of these breast asymmetry  
335 measures as diagnostic or predictor markers will require the conduct of large-scale and longitudinal  
336 studies with objective measurements of breast asymmetry. Objective breast tissue asymmetry  
337 estimates can now be obtained using existing fully-automated mammographic volumetric analysis  
338 tools and thus can be provided, without additional investigations, for all women attending screening.  
339 The availability of such data will facilitate further research into the association between asymmetry  
340 and breast cancer, both at the current screen and subsequently, and may potentially provide a  
341 practical additional tool for stratifying the screening population in terms of likelihood of having, or  
342 risk of developing, breast cancer.

#### 343 **LIST OF ABBREVIATIONS**

344 BV: breast volume; CC: cranio-caudal view; SWLBSS: South West London breast screening service;  
345 DV: dense volume i.e. absolute volume of dense tissue; IMD Index of Multiple Deprivation; IQR:  
346 Interquartile Range; MLO: mediolateral oblique mammogram view; %MD: percentage  
347 mammographic breast density; MRI: magnetic resonance imaging; NHS: National Health Service;  
348 NHSBSP: England and Wales NHS Breast Screening Programme; SD: standard deviation; Volpara:  
349 Volpara density algorithm.

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