

# Cloudiness and Breast Cancer

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## Abstract

Traditional risk factors for breast cancer explain only a fraction of cases. Causes for trends in breast cancer incidence are not fully understood. Breast cancer incidence and mortality rates decrease with environmental conditions that promote Vitamin D synthesis in human skin including lower latitude and higher personal exposure to sunlight. Association of temporal variability in breast cancer incidence with changes in cloudiness, which decrease human Vitamin D synthesis is investigated. Association between temporal changes in breast cancer incidence and in the autumn cloudiness for preceding years is computed using data for the United States. There is a correlation of 0.96 (95%CI=0.92-1) between the time series of breast cancer incidence in the age group of 70-79 years and the average cloudiness in October during preceding 20 years. An empirical model for breast cancer incidence using autumn cloudiness in preceding years captures a rapid increase in breast cancer incidence in the 1980s and some of its year-to-year variability. Increased autumn cloudiness is associated with increased subsequent breast cancer incidence. Proposed mechanism includes blocking of solar ultraviolet radiation by thick clouds and decreasing the synthesis of Vitamin D in human skin. The findings suggest a new connection between climate variability and human health.

**Keywords:** Breast neoplasms; Climate; Incidence; SEER Program; Ultraviolet rays; United States; Vitamin D

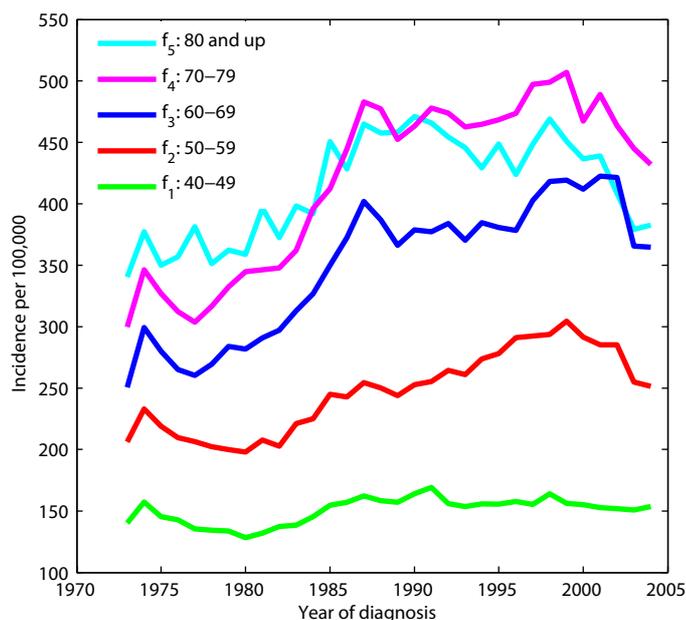
**Abbreviations:** HRT: Hormone Replacement Therapy; EPRT: Estrogen-plus-Progestin Replacement Therapy; ER+: Estrogen Receptor positive; ISCCP: International Satellite Cloud Climatology Project; SEER: Surveillance, Epidemiology and End Results; US: United States; UV: Ultraviolet; WMO: World Meteorological Organization

## Introduction

Breast cancer incidence rates for several representative US regions have been recorded by the Surveillance, Epidemiology, and End Results (SEER) Program of the National Cancer Institute (SEER, 2007) since 1973. Time series of age-adjusted incidence rates  $f_1(t), f_2(t), \dots, f_5(t)$  for five age groups of women are shown in Figure 1. The incidence rates for women diagnosed between 70 and 79 years of age ( $f_4(t)$ : magenta curve) climbed rapidly from mid 1970s to mid 1980s and declined from 1999 to 2004. The introduction of screening mammography, which allows earlier detection of cancer, contributed to the former increase (Miller et al., 1993), but it cannot account for the entire observed increase (White et al., 1990). A decline in the use of hormone replacement therapy (HRT) (Glass et al., 2007) or saturation in screening mammography (Li and Darling, 2007) may

have contributed to the latter decline in cancer incidence. Age, early menarche, nulliparity, late age of first full-term pregnancy, late age and short length of lactation, late menopause, genetic mutations, use of HRT, radiation exposure, alcohol consumption, higher educational level and higher socioeconomic status are among risk factors for breast cancer, many of which are related to cumulative exposure to estrogens (Coyle, 2004). However, causes for trends and year-to-year variations in breast cancer incidence are not fully understood (Garfinkel, 1993; Ghafoor et al., 2003).

Low vitamin D status is a risk factor for the development of breast cancer (Coyle, 2004; John et al., 1999). Synthesis in the skin exposed to solar UV radiation is a major source of Vitamin D in humans (Holick, 2004). Breast cancer incidence and mortality are higher in geographic regions with lower UV radiation (Gorham et al., 1990; Garland et al., 1990; Grant, 2002; Grant and Garland, 2006). In the extratropics low wintertime UV amounts may be



**Figure 1:** Breast cancer incidence. Annual rates per 100,000 females of any race are shown by the age at diagnosis:  $f_1$  (age 40-49 years),  $f_2$  (age 50-59),  $f_3$  (age 60-69),  $f_4$  (age 70-79) and  $f_5$  (age 80 and above). Age adjusted incidence rates to standard US population in year 2000 are shown (SEER, 2007). Statistics were generated from malignant cases only.

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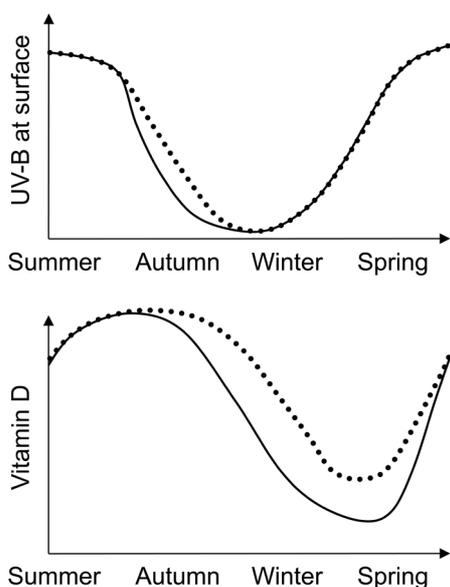
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inadequate for vitamin D synthesis (Webb et al., 1988) causing seasonal variability in serum Vitamin D status (Lappe et al., 2006; Kull et al., 2009). Due to Vitamin D half-life of 1-2 months (Vieth, 1999), a deficiency is common at the end of the winter (Holick, 2004). Assuming that autumn vitamin D synthesis modulates subsequent Vitamin D minima (Figure 2), we ask: Is interannual variability in the autumn surface UV radiation related to breast cancer incidence?

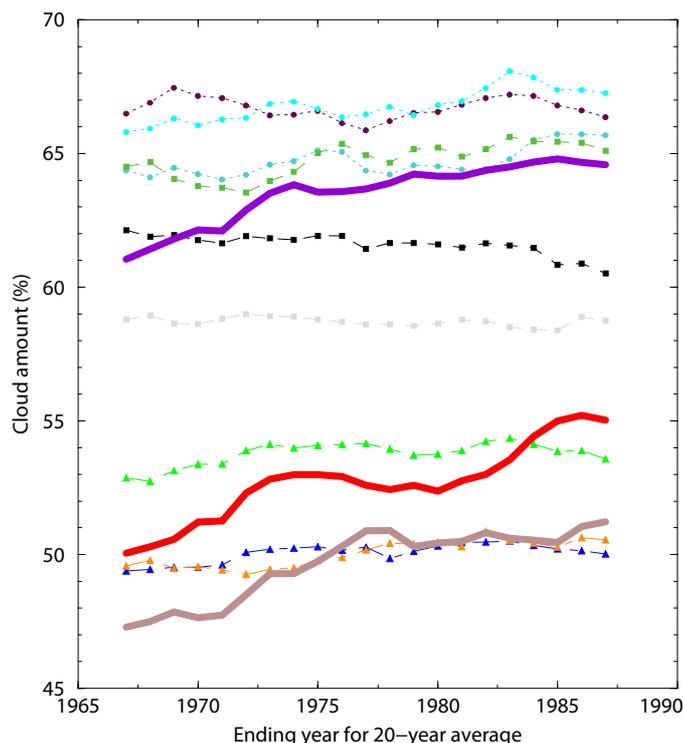
High quality, multi-decadal observations of surface UV are needed to examine temporal association between UV radiation and breast cancer incidence. However, ground-based observations providing time series since 1960s are of insufficient quality to establish trends (Weatherhead et al., 1997). A newer network of ground based stations has been collecting UVB observations since 1990s (Bigelow et al., 1998), but their record is too short at the time of this writing. Assuming that exposure is relevant over about two decades from initiation to detection of breast cancer, about 50 years of UV data are needed to evaluate association with the 30 years of subsequent SEER breast cancer incidence rates. Satellite estimates of surface UV have difficulties over polluted sites (McKenzie et al., 2001), where they do not capture reduction in UV radiation caused by urban aerosols near the ground (Herman et al., 1999). Solar UV radiation is scattered and absorbed in the atmosphere by clouds, ozone, aerosols, and nitrogen dioxide (WMO, 2003). Thick clouds can reduce UV-B radiation at wavelengths of 290-320 nm by 80% (Frederick and Steele, 1995). The focus of this study is on interannual variability in cloudiness due to its large impact on surface UV.

**Materials and Methods**

For 1987 and earlier years we use monthly mean daylight cloud amounts from an edited record of all US ground stations (Karl and Steurer, 1990), which was used to study cloudiness trends over the United States. Due to changes in observing practices spurious trends may exist between late 1930s and late 1940s. An increasing trend in cloudiness and a consistent decrease in



**Figure 2:** Schematic annual cycle. Idealized annual cycle is shown for a) UV-B at the surface for cloud-free conditions (dotted) and for cloudy autumn (solid) and for b) corresponding serum 25-hydroxyvitamin D [25(OH)D] concentration.

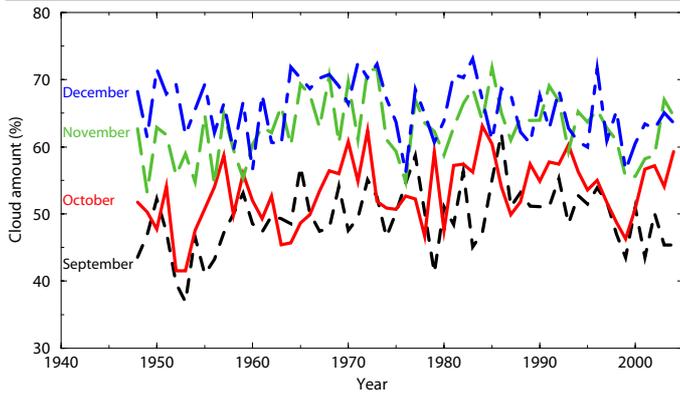


**Figure 3:** Average cloudiness for each month for the preceding 20 years. Months are marked by season: winter (circles), spring (squares) and summer (triangles). Autumn months are shown by thicker lines: September (brown), October (red) and November (violet).

temperature range were found since 1948. From each of the nine SEER regions we selected one ground station with complete records from 1948 to 1987. Two stations are available in Iowa; Sioux City is very close to the state border, so Des Moines was selected to represent this state. Only one station has a complete record in each of the other four SEER states: Hartford in Connecticut, Albuquerque in New Mexico, Salt Lake City in Utah, and Honolulu in Hawaii. For each of the four SEER metropolitan areas (San Francisco, California; Seattle, Washington; Detroit, Michigan; and Atlanta, Georgia) there is a ground station for that metropolitan area.

Variability in cloudiness over several years is examined using the spatial average over the nine stations in SEER regions. For each month this cloudiness was averaged for 20-year intervals. From the 20-year interval ending in 1967 to the 20-year interval ending in 1987 the largest increase in cloudiness is seen for October (5%), followed by September (4%), and November (3.5%, Figure 3). Due to this substantial increase in cloudiness in the autumn months, we focus the remaining analysis on the autumn months.

Monthly cloudiness records for the nine stations in SEER regions were extended to year 2004 using monthly total cloud amounts from satellite based International Satellite Cloud Climatology Project (ISCCP) data (Rossow and Schiffer, 1999). Measurements from geostationary and polar orbiting satellites are combined every 3 hours and subsequently averaged to produce ISCCP monthly total cloud amounts. Multiple satellites provide measurements over some locations. Thus, ISCCP uses a hierarchy of preferred sources of satellite data that based on satellite observing geometries, with preference for data from the satellite



**Figure 4:** Monthly average cloudiness. Average cloud amounts for the nine ground stations in SEER regions are shown for September ( $g_j$ ; black), October ( $g_{10}$ ; red), November ( $g_{11}$ ; green) and December ( $g_{12}$ ; blue).

closest to being overhead rather than low on the horizon. For the 48 contiguous United States and Hawaii the primary data sources are geostationary satellites (Geostationary Operational Environmental Satellite (GOES)-East and GOES-West). The monthly maps of cloudiness from ISCCP allow us to examine how well cloudiness over chosen stations represents cloudiness over SEER regions. For example, the October cloudiness from 1983 to 2004 over Des Moines and the cloudiness over a rectangular area representing Iowa both vary from <60% to >75%; even though cloudiness for Des Moines is 1.8% lower, there is a correlation of 0.95 between their time series. Differences between ground-based and satellite observations during the 1980-1987 overlap period are typically smaller than the interannual variability in the records at each of the nine stations. Figure 4 shows the time series of the nine-station average of the cloud amounts for September, October, November, and December, which are denoted by  $g_9, g_{10}, g_{11}$ , and  $g_{12}$ , respectively.

In order to examine the relationship between multi-annual averages of cloudiness in one of the autumn months and the breast cancer incidence, we formulate the following model  $h$  to compare with breast cancer incidence  $f_i$ :

$$h(t) = c + \frac{b}{l} \sum_{k=a}^{a+l-1} g_j(t-k), \text{ for } j = 9, 10, 11, 12.$$

In this model monthly cloud amounts  $g_j$  are averaged over  $l$  years, with a lag of  $a$  years. In order to match ranges of  $f_i$  and  $h$ , linear stretching is controlled by parameter  $b$  and offsets by  $c$ .

For example, for  $j=10$ ,  $g_j$  is cloudiness for October. Cloudiness is averaged over several ( $l$ ) years because cancer may be influenced by conditions that occurred over decades before its detection. The lag parameter  $a$  is introduced to allow a possibility that cloudiness in the last few years may be irrelevant, e.g. if the cancer is too advanced to be influenced by saddle changes in Vitamin D sources. All cloudiness attains values between 0 (clear sky) and 100 (cloud covered sky), so by construction, averages of cloudiness will produce numbers between 0 and 100. In order to fit breast cancer incidence (which e.g. varies from ~300/100,000 to ~500/100,000 for women in the age group 70-79 as seen in Figure 1) parameter  $b$  allows adjusting the range of variability, and the parameter  $c$  allows vertical offsets to bring curves closer together.

Using least-squares norm we determine the best fit  $h_i(t)$  to  $f_i(t)$ , for  $i=1,2,\dots,5, j=9,10,11,12, a \geq 0, l \geq 1$  and  $a+l \leq 25$ . The causality limits  $a$  to be positive. Total temporal shifts ( $a+l$ ) do not exceed 25 years to avoid artificial offsets in cloudiness data in 1930s and 1940s caused by changes in cloud measurement techniques (Karl and Steurer, 1990).

**Results**

For the age group 70-79 the best least-squares fit to  $f_4$  is provided by  $h_4$  using cloudiness in October ( $j=10$ ) and the following parameters:

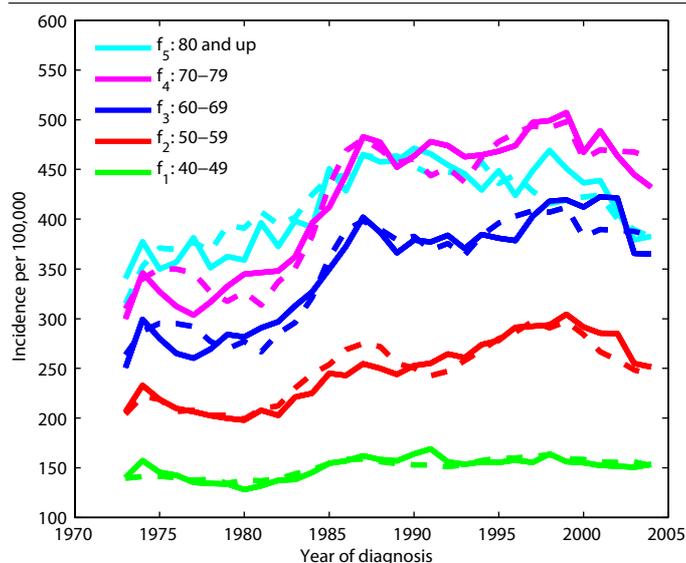
$$a = 1, l = 20, b = 58.78, \text{ and } c = -2764.9.$$

Thus, the best fit uses October cloudiness for 1 to 20 ( $a$  to  $a+l-1$ ) years before the cancer incidence. Note the importance of parameter  $b$ : it quantifies the increase of cancer incidence rate per 100,000 individuals corresponding to a 1% increase in the average cloudiness over the  $l$  years in the model  $h$ . Cloudiness for October or November provided best fit for all five age groups, with the parameters in Table 1. The correlation coefficients between the incidence and best fit models range from 0.84 (95%CI=0.70-0.95) to 0.96 (95%CI=0.92-1). The time series of best fit models in Figure 5 capture the strong increases in breast cancer incidence rates during 1980s and more stable rates since 1987.

Spatial association between cloudiness and breast cancer was investigated as well. There is a correlation of 0.815 between 20-year average (1983-2002) of October cloudiness (Rossow and Schiffer, 1999) over regions included in the SEER program and the breast cancer incidence in 2003 for white women older than

Age group	Group index $i$	Month $j$	Number of years $l$	Lag $a$	Sensitivity to change In Cloudiness $b$	Offset $c$	Correlation between $f_i$ and $h_i$	95% confidence interval
40-49	1	10	19	1	7.9513	-280.8	0.8408	0.6962-0.9537
50-59	2	10	17	2	26.3441	-1182.2	0.9342	0.8684-0.9924
60-69	3	10	20	1	46.3609	-2151.3	0.9483	0.8958-0.9953
70-79	4	10	20	1	58.7847	-2764.9	0.9599	0.9187-0.9972
80+	5	11	23	2	47.8014	-2636.2	0.8704	0.7490-0.9698

**Table 1:** The parameters for model  $h_i$  that provide best fit using least square norm to the breast cancer incidence  $f_i$  for five age groups.



**Figure 5:** Empirical model using best fit parameters. Breast cancer incidence  $f_1$  to  $f_5$  for women of ages 40-49, 50-59, 60-69, 70-79, 80 and older (solid: colors as in Figure 1) and the fit functions  $h_1$  to  $h_5$  (dashed) with parameters given in Table 1.

65 years (Table 2). Note that solar zenith angle, cloudiness and ozone, which all decrease surface UV-B, typically increase with increasing latitude, so the correlation is not likely to be due to changes in cloudiness alone. Latitude of SEER cities and geographic centers for SEER states is also given in Table 2. There is a negative correlation of -0.366 between latitude and breast cancer incidence rates, indicating that latitude alone does not explain variability in breast cancer incidence rates. The surprising negative correlation is due to Hawaii data. When correlations are computed for 8 SEER regions, excluding Hawaii, there is a positive correlation of 0.564 between latitude and breast cancer incidence. Correlation between cloudiness and breast cancer is much less sensitive to exclusion of Hawaii data; it drops only slightly to 0.813.

**Discussion**

High temporal correlations between autumn cloudiness and breast cancer incidence exist, which do not imply causality. However, we assert that synthesis of Vitamin D in human skin is the linking mechanism, which is consistent with earlier studies of environmental exposure to solar radiation (John et al., 1999;

Gorham et al., 1990; Garland et al., 1990; Grant, 2002). Dietary supplementation provides an alternative source of Vitamin D and another mode of intervention in order to reduce the burden of breast cancer. Randomized trials of daily Vitamin D and calcium supplementation in postmenopausal women showed mixed results: 400 IU of Vitamin D and 1000 mg of elemental calcium did not reduce colon cancer incidence (Wactawski-Wende et al., 2006), but 1100 IU of Vitamin D plus 1400-1500 mg of calcium substantially reduced incidence of breast, colon, and several other cancers (Lappe et al., 2007). The latter trial, which is consistent with our findings, had a smaller sample population, but its strengths lie in the higher dose of Vitamin D and a better adherence to treatment.

This is the first study into factors influencing autumn synthesis of Vitamin D and subsequent cancer incidence. Several factors modulating surface UV radiation in the autumn may be important. This study focuses on the large variability caused by clouds. Spatial correlation of 0.815 was found between autumn cloudiness and subsequent breast cancer incidence, which is much stronger than the spatial correlation that was found between latitude and breast cancer incidence. Correlations between breast cancer incidence and annual mean of the total solar radiation (-0.75, Gorham et al., 1990) are weaker. Correlations between breast cancer mortality and annual mean of the total solar radiation (-0.80, Garland et al., 1990) or summertime UV radiation (-0.67, Grant, 2002) are weaker as well. Wintertime UV radiation may be inadequate for synthesis of Vitamin D (Webb et al., 1988), making autumn the key period for synthesis of Vitamin D that is used during winter. In contrast, cloudiness in the spring is not expected to have a lasting effect because UV radiation is generally increasing during this season.

High temporal correlations of up to 0.96 were found between cloudiness in the autumn months, especially October, and subsequent breast cancer incidence. This temporal correlation is stronger than any of the previously discussed spatial correlations. Variability in cloudiness is a major factor causing interannual variability in UV radiation at the surface. Other factors include variability in ozone, aerosols or nitrogen dioxide (WMO, 2003). Interannual ozone variability in the northern middle latitudes is the lowest in the autumn (September to November) among all the seasons (WMO, 2003). Changes in atmospheric aerosol concentrations may have caused a 5% decline in the solar radiation

	New Mexico	Utah	Atlanta	Connecticut	Seattle	Detroit	Iowa	San Francisco	Hawaii
<b>Cloudiness in October 1983-2002 [%]</b>	38.8	42.65	50.55	56.425	59.925	60.875	61	64.075	66.225
<b>Latitude [degrees N]</b>	34.50	39.38	33.65	41.60	47.45	42.42	41.96	37.62	20.95
<b>Breast cancer incidence rate per 100,000 for 65+</b>	368.4	374.4	413.2	419.7	490.5	430.4	409.2	472.4	542.5

**Table 2:** Average cloudiness, latitude (of geographic center for states), and breast cancer incidence for white females older than 65 years in 9 SEER regions.

incident at the Earth's surface from 1960 to 1990 and a later increase through 1990s (Wild et al., 2005). Aerosol variability could also be related to breast cancer incidence, and is a topic for future research.

Stratification of cloudiness-cancer correlations by age group or time period provides insight into possible roles of Vitamin D. First, among all age groups correlations are the highest for ages 70-79, when occurrence of estrogen receptor positive (ER+) breast cancer peaks (Anderson and Matsuno, 2006). Given that temporal variability of less common estrogen receptor negative cancer differs (Glass et al., 2007) from that of ER+, this suggests a stronger link between cloudiness and ER+ breast cancer. Second, restricting the analysis to periods receding widespread estrogen-plusprogesterin replacement therapy (EPRT), the highest correlation of 0.972 is found between 1973-1988 breast cancer incidence for ages 70-79 and cloudiness for 12 Decembers at least 13 years before the cancer detection. This suggests a preventative role of vitamin D in initial phases of cancerogenesis. Third, inclusion of data for 1990s shifts best fit to October cloudiness. Estrogen therapy alone raises levels of the highly active 1,25-dihydroxyvitamin D, but the addition on progesterin lowers its levels (Bikkle et al., 1992). Vitamin D requirements might increase during EPRT shifting the sensitivity to October cloudiness, when UV radiation is stronger, and to the years closer to the cancer detection, when vitamin D may play a role in suppressing the growth of still undetected cancer that is fueled by the EPRT (Dietel et al., 2005).

Since the release of Women's Health Initiative trial results, the use of EPRT has declined substantially. Ravdin et al., (2007) attributed the subsequent reduction of breast cancer incidence rates to the decline in the use of EPRT. However, it is plausible that vitamin D supplementation increased concurrently and also contributed to the reduction of breast cancer incidence rates. Such increase in vitamin D supplementation would be consistent with recent trends that show almost five-fold increase in vitamin D prescriptions from year 2006 to year 2008 (Drug Topics, 2009). Additional data are needed on recent trends in serum vitamin D levels. Note that a marked decrease in serum 25(OH) D levels was observed from the 1988-1994 to the 2001-2004 National Health and Nutrition Examination Survey (NHANES) data collections (Ginde et al., 2009).

It is recognized that factors other than ambient UV light exposure, such as personal behavior, melanin content, and age, have impact on Vitamin D synthesis and Vitamin D status (Holick, 2004; McCarty, 2008). Avoidance of UV radiation in order to decrease risk of skin cancer is common and it may be leading to Vitamin D deficiency. For example, sunscreen efficiency and use have increased. Under very controlled conditions the sunscreen can completely suppress cutaneous Vitamin D synthesis. However, when subjects were allowed to apply their own sunscreen no significant changes in serum concentrations of Vitamin D were found (McCarty, 2008). A limitation of the present study is that associations were observed in groups and may not apply to individuals within these groups. Thus, further study of individual exposure, serum vitamin D levels and breast cancer risk is needed. Higher personal exposure to sunlight determined by physician assessment or self-reported frequent recreational or occupational exposure were shown to lower breast cancer risk (John et al., 1999). Analysis of personal skin pigmentation mea-

surements showed a reduced risk of advanced breast cancer among women with naturally light skin pigmentation and higher pigmentation on the forehead, which indicates higher exposure to sunlight (John et al., 2007). Furthermore, several recent case-control studies found an inverse association between serum vitamin D levels and breast cancer risk (Blackmore et al., 2008; Abbas et al., 2008; Crew et al., 2009; Rossi et al., 2009). A future study that takes measurements of Vitamin D status at the end of the winter following varying autumn cloudiness over several years could provide further information about the asserted connection among cloudiness and Vitamin D status, and potential links to breast cancer incidence.

High temporal correlations between autumn cloudiness and subsequent breast cancer incidence offer an alternative to previously proposed explanations of breast cancer trends (Miller et al., 1993; White et al., 1990; Glass et al., 2007; Li and Darling, 2007; Garfinkel, 1993; Ghafoor et al., 2003). The length of breast cancer incidence time series is a limitation of the study. Its strengths lie in the use of independently collected datasets, focus on the autumn season, and on cloudiness as a major modulator of the solar UV reaching the Earth's surface. The proposed mechanism of action through modulation of Vitamin D synthesis in human skin provides a new example of possible interaction between climate variability and human health on decadal scales.

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### References

1. Abbas S, Linseisen J, Slanger T, Kropp S, Mutschelknauss EJ, et al. (2008) Serum 25-hydroxyvitamin D and risk of post-menopausal breast cancer—results of a large case-control study. *Carcinogenesis* 29: 93-99. » [CrossRef](#) » [PubMed](#) » [Google Scholar](#)
2. Anderson WF, Matsuno R (2006) Breast cancer heterogeneity: a mixture of at least two main types. *J Natl Cancer Inst* 98: 948-51. » [CrossRef](#) » [PubMed](#) » [Google Scholar](#)
3. Bigelow DS, Slusser JR, Beaubien AF, Gibson JH (1998) The USDA Ultraviolet Radiation Monitoring Program. *Bull Amer Meteorol Soc* 79: 601-615. » [CrossRef](#) » [Google Scholar](#)
4. Bikle DD, Halloran BP, Harris ST, Portale AA (1992) Progesterin antagonism of estrogen stimulated 1,25-dihydroxyvitamin D levels. *J Clin Endocrinol Metab* 75: 519-23. » [CrossRef](#) » [PubMed](#) » [Google Scholar](#)
5. Blackmore KM, Lesosky M, Barnett H, Raboud JM, Vieth R, et al. (2008) Vitamin D From Dietary Intake and Sunlight Exposure and the Risk of Hormone-Receptor-Defined Breast Cancer. *Am J Epidemiol* 168: 915-924. » [CrossRef](#) » [PubMed](#) » [Google Scholar](#)
6. Coyle YM (2004) The effect of environment on breast cancer risk. *Breast Cancer Res Treat* 84: 273-288. » [CrossRef](#) » [PubMed](#) » [Google Scholar](#)
7. Crew KD, Gammon MD, Steck SE, Hershman DL, Cremers S, et al. (2009) Association between plasma 25-hydroxyvitamin D and breast cancer risk. *Cancer Prev Res* 2: 598-604. » [CrossRef](#) » [PubMed](#) » [Google Scholar](#)
8. Dietel M, Lewis MA, Shapiro S (2005) Hormone replacement therapy: pathobiological aspects of hormone-sensitive cancers in women relevant to epidemiological studies on HRT: a mini-review. *Hum Reprod* 20: 2052-60. » [CrossRef](#) » [PubMed](#) » [Google Scholar](#)
9. Drug Topics (2009) Drugs by units in the United States in specific years.
10. Frederick JE, Steele HD (1995) The transmission of sunlight through

- cloudy skies: An analysis based on standard meteorological information. *J Appl Meteorol* 34: 2755-2761. » [CrossRef](#) » [Google Scholar](#)
11. Garfinkel L (1993) Current trends in breast cancer. *CA Cancer J Clin* 43: 5-6. » [CrossRef](#) » [PubMed](#) » [Google Scholar](#)
  12. Garland FC, Garland CF, Gorham ED, Young JF (1990) Geographic variation in breast cancer mortality in the United States: a hypothesis involving exposure to solar radiation. *Prev Med* 19: 614-622. » [PubMed](#) » [Google Scholar](#)
  13. Ghafoor A, Jemal A, Ward E, Cokkinides V, Smith R, et al. (2003) Trends in Breast Cancer by Race and Ethnicity. *CA Cancer J Clin* 53: 342-355. » [CrossRef](#) » [PubMed](#) » [Google Scholar](#)
  14. Ginde AA, Liu MC, Camargo CA Jr (2009) Demographic Differences and Trends of Vitamin D Insufficiency in the US Population, 1988-2004. *Arch Intern Med* 169: 626-632. » [CrossRef](#) » [PubMed](#) » [Google Scholar](#)
  15. Glass AG, Lacey JV Jr, Carreon JD, Hoover RN (2007) Breast cancer incidence, 1980-2006: combined roles of menopausal hormone therapy, screening mammography, and estrogen receptor status. *J Natl Cancer Inst* 99: 1152-61. » [CrossRef](#) » [PubMed](#) » [Google Scholar](#)
  16. Gorham ED, Garland FC, Garland CF (1990) Sunlight and breast cancer incidence in the USSR. *Int J Epidemiol* 19: 820-824. » [CrossRef](#) » [PubMed](#) » [Google Scholar](#)
  17. Grant WB (2002) An estimate of premature cancer mortality in the U.S. due to inadequate doses of solar ultraviolet-B radiation. *Cancer* 94: 1867-75. » [CrossRef](#) » [PubMed](#) » [Google Scholar](#)
  18. Grant WB, Garland CF (2006) The association of solar ultraviolet B (UVB) with reducing risk of cancer: multifactorial ecologic analysis of geographic variation in age-adjusted cancer mortality rates. *Anticancer Res* 26: 2687-99. » [CrossRef](#) » [PubMed](#) » [Google Scholar](#)
  19. Herman JR, Krotkov N, Celarier E, Larko D, Labow G (1999) Distribution of UV radiation at the Earth's surface from TOMS-measured UV-backscattered radiances. *J Geophys Res* 104: 12059-12076. » [CrossRef](#) » [Google Scholar](#)
  20. Holick MF (2004) Sunlight and vitamin D for bone health and prevention of autoimmune diseases, cancers and cardiovascular disease. *Am J Clin Nutr* 80: 1678S-88S. » [CrossRef](#) » [PubMed](#) » [Google Scholar](#)
  21. John EM, Schwartz GG, Dreon DM, Koo J (1999) Vitamin D and Breast Cancer Risk: The NHANES I Epidemiologic Follow-up Study, 1971-1975 to 1992. *Cancer Epidemiology Biomarkers & Prevention* 8: 399-406. » [CrossRef](#) » [PubMed](#) » [Google Scholar](#)
  22. John EM, Schwartz GG, Koo J, Wang W, Ingles SA (2007) Sun exposure, vitamin D receptor gene polymorphisms, and breast cancer risk in a multiethnic population. *Am J Epidemiol* 166: 1409-19. » [CrossRef](#) » [PubMed](#) » [Google Scholar](#)
  23. Karl TR, Steurer PM (1990) Increased cloudiness in the United States during the first half of the twentieth century: Fact or fiction. *Geophys Res Lett* 17: 1925-1928. » [CrossRef](#) » [Google Scholar](#)
  24. Kull M, Kallikorm R, Tamm A, Lember M (2009) Seasonal variance of 25-(OH) vitamin D in the general population of Estonia, a Northern European country. *BMC Public Health* 9: 22. » [CrossRef](#) » [PubMed](#) » [Google Scholar](#)
  25. Lappe JM, Davies KM, Travers-Gustafson D, Heaney RP (2006) Vitamin D status in a rural postmenopausal female population. *J Am Coll Nutr* 25: 395-402. » [CrossRef](#) » [PubMed](#) » [Google Scholar](#)
  26. Lappe JM, Travers-Gustafson D, Davies KM, Recker RR, Heaney RP (2007) Vitamin D and calcium supplementation reduces cancer risk: results of a randomized trial. *Am J Clin Nutr* 85: 1586-91. » [CrossRef](#) » [PubMed](#) » [Google Scholar](#)
  27. Li CI, Darling JR (2007) Changes in Breast Cancer Incidence Rates in the United States by Histologic Subtype and Race/Ethnicity, 1995 to 2004. *Cancer Epidemiol Biomarkers Prev* 16: 2773-2780. » [CrossRef](#) » [PubMed](#) » [Google Scholar](#)
  28. McCarty CA (2008) Sunlight exposure assessment: can we accurately assess vitamin D exposure from sunlight questionnaires. *Am J Clin Nutr* 87: 1097S-1101S. » [CrossRef](#) » [PubMed](#) » [Google Scholar](#)
  29. McKenzie RL, Seckmeyer G, Bais AF, Kerr JB, Madronich S (2001) Satellite retrievals of erythemal UV dose compared with ground-based measurements at northern and southern midlatitudes. *J Geophys Res* 106: 24051-24062. » [CrossRef](#) » [Google Scholar](#)
  30. Miller BA, Feuer EJ, Hankey BF (1993) Recent incidence trends for breast cancer in women: An update. *CA Cancer J Clin* 43: 27-41. » [CrossRef](#) » [PubMed](#) » [Google Scholar](#)
  31. Ravdin PM, Cronin KA, Howlander N, Berg CD, Chlebowski RT, et al. (2007) The decrease in breast-cancer incidence in 2003 in the United States. *Engl J Med* 356: 1670-4. » [CrossRef](#) » [PubMed](#) » [Google Scholar](#)
  32. Rossi M, McLaughlin JK, Lagiou P, Bosetti C, Talamini R, et al. (2009) Vitamin D intake and breast cancer risk: a case-control study in Italy. *Ann Oncol* 20: 374-8. » [CrossRef](#) » [PubMed](#) » [Google Scholar](#)
  33. Rossow WB, Schiffer RA (1999) Advances in understanding clouds from ISCCP. *Bull of the Amer Meteor Soc* 80: 2261-2287. » [CrossRef](#) » [Google Scholar](#)
  34. Surveillance, Epidemiology, and End Results (SEER) Program ([www.seer.cancer.gov](http://www.seer.cancer.gov)) SEER\*Stat Database: Incidence - SEER 9 Regs Limited-Use, Nov 2006 Sub (1973-2004), National Cancer Institute, DCCPS, Surveillance Research Program, Cancer Statistics Branch, released April 2007, based on the November 2006 submission.
  35. Vieth R (1999) Vitamin D supplementation, 25-hydroxyvitamin D concentrations, and safety. *Am J Clin Nutr* 69: 842-56. » [CrossRef](#) » [PubMed](#) » [Google Scholar](#)
  36. Wactawski-Wende J, Kotchen JM, Anderson GL, Assaf AR, Brunner RL, et al. (2006) Calcium plus vitamin D supplementation and the risk of colorectal cancer. *N Engl J Med* 354: 684-96. » [CrossRef](#) » [PubMed](#) » [Google Scholar](#)
  37. Weatherhead EC, Taio GC, Reinsel GC, Frederick JE, DeLuisi JJ, et al. (1997) Analysis of long-term behavior of ultraviolet radiation measured by Robertson-Berger meters at 14 sites in the United States. *J Geophys Res* 102: 8737-8754. » [CrossRef](#) » [Google Scholar](#)
  38. Webb AR, Kline L, Holick MF (1988) Influence of season and latitude on the cutaneous synthesis of vitamin D3: exposure to winter sunlight in Boston and Edmonton will not promote vitamin D3 synthesis in human skin. *J Clin Endocrinol Metab* 67: 373-8. » [CrossRef](#) » [PubMed](#) » [Google Scholar](#)
  39. White EW, Lee CY, Kristal AR (1990) Evaluation of the increase in breast cancer incidence in relation to mammography use. *J Natl Cancer Inst* 82: 1546-1552. » [CrossRef](#) » [PubMed](#) » [Google Scholar](#)
  40. Wild M, Gilgen H, Roesch A, Ohmura A, Long CN, et al. (2005) From Dimming to Brightening: Decadal Changes in Solar Radiation at Earth's Surface. *Science* 308: 847-850. » [CrossRef](#) » [PubMed](#) » [Google Scholar](#)
  41. World Meteorological Organization (2003) Scientific assessment of ozone depletion: 2002, in *Global Ozone Research and Monitoring Project*.

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