**Name of Journal: *World Journal of Orthopedics***

**ESPS Manuscript NO: 24189**

**Manuscript Type: Original Article**

***Retrospective Study***

**Seasonal variation in adult hip disease secondary to osteoarthritis and developmental dysplasia of the hip**

Sueyoshi T *et al*. Seasonal variation of OA and DDH

**Tatsuya Sueyoshi, Merrill A Ritter, Kenneth E Davis, Randall T Loder**

**Tatsuya Sueyoshi,** Department of Orthopedic Surgery, Kyoto University, Kyoto 606-8507, Japan

**Merrill A Ritter, Kenneth E Davis,** Center for Hip and Knee Surgery, Mooresville, IN 46158, United States

**Randall T Loder,** James Whitcomb Riley Children’s Hospital, Indianapolis, IN 46202, United States

**Author contributions:** Sueyoshi T primary contributed to the work and wrote the manuscript; Ritter MA did the surgery and organized the study; Davis KE did the data collection and statistics; Loder RT made some of DDH comments and gave suggestion.

**Institutional review board statement:** The study was reviewed and approved by the St. Francis Hospital Institutional Review Board.

**Informed consent statement:** All study participants or their legal guardian provided consent about personal and medical data collection prior to study enrolment.

**Conflict-of-interest statement:** All the Authors have no conflict of interest related to the manuscript.

**Data sharing statement:** The original anonymous dataset is available on request from the corresponding author at jrsiresearch1@gmail.com.

**Open-Access:** This article is an open-access article which was selected by an in-house editor and fully peer-reviewed by external reviewers. It is distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited and the use is non-commercial. See: <http://creativecommons.org/licenses/by-nc/4.0/>

**Manuscript source:** Unsolicited manuscript

**Corresponding to: Merrill A Ritter, MD,** Center for Hip and Knee Surgery, 1199 Hadley Road, Mooresville, IN 46158, United States. jrsiresearch1@gmail.com

**Telephone:** +1-317-8345263

**Received:** January 12, 2016

**Peer-review started:** January 14, 2016

**First decision:** March 7, 2016

**Revised:** August 8, 2016

**Accepted:** August 17, 2016

**Article in press:**

**Published online:**

**Abstract**

**AIM:** To determine if there was a seasonal variation in adults undergoing total hip arthroplasty for end stage hip disease due to osteoarthritis (OA) or sequelae of developmental dysplasia of the hip (DDH).

**METHODS:** The total hip registry from the author’s institution for the years 1969 to 2013 was reviewed. The month of birth, age, gender, and ethnicity was recorded. Differences between number of births observed and expected in the winter months (October through February) and non-winter mo (March through September) were analyzed with the *χ*2 test. Detailed temporal variation was mathematically assessed using cosinor analysis.

**RESULTS:** There were 7792 OA patients and 60 DDH patients who underwent total hip arthroplasty. There were more births than expected in the winter months for both the DDH (*P* < 0.0001) and OA (*P* = 0.0052) groups. Cosinor analyses demonstrated a peak date of birth on October 1st.

**CONCLUSION:** These data demonstrate an increased prevalence of DDH and OA in those patients born in winter.

**Key words:** Seasonal trend; Winter; Birth month; Developmental hip dysplasia; Osteoarthritis

**© The Author(s) 2016.** Published by Baishideng Publishing Group Inc. All rights reserved.

**Core tip:** The purpose of this study was to determine if there was a seasonal variation undergoing total hip arthroplasty for osteoarthritis (OA) or developmental dysplasia of the hip (DDH). Differences between number of births observed and expected in the winter months and non-winter months were analyzed with the *χ*2 test. There were 7792 OA and 60 DDH, and more births than expected in the winter months for both the DDH (*P* < 0.0001) and the OA (*P* = 0.0052) cohorts. These data clearly demonstrated an increased prevalence of DDH and OA in those patients born during the winter months.

Sueyoshi T, Ritter MA, Davis KE, Loder RT. Seasonal variation in adult hip disease secondary to osteoarthritis and developmental dysplasia of the hip. *World J Orthop* 2016; In press

**INTRODUCTION**

Total hip arthroplasty (THA) is a standard, successful treatment for end stage hip disease. The etiology of such disease is frequently due to osteoarthritis (OA), osteonecrosis/avascular necrosis (ON/AVN), rheumatoid arthritis (RA) and developmental dysplasia of the hip (DDH). Studies of hip osteoarthritis suggest that childhood deformities or residuals of childhood hip disease often experience accelerated or premature joint degeneration[1,2]. DDH is a wide spectrum of deformity ranging from mild acetabular dysplasia to complete and fixed dislocation[3]. Many studies have noted that DDH is more frequent in patients born in the colder months[4-6]. DDH with residual acetabular dysplasia is often followed by OA of the hip. OA has also been reported to be more frequent in people born in the winter[7]. The authors wished to determine if there was any seasonal variation in adults who are treated for OA and DDH in the Midwest region of the United States. This will further augment a previous study which noted a seasonal variation in children with DDH in this region[3].

**MATERIALS AND METHODS**

A retrospective review of all THAs performed from November 1969 through November 2013 at the author’s institution was performed. This review was approved by the local Institutional Review Board. There were 10572 THAs in 8579 patients that met the initial inclusion criteria of primary OA, RA, ON or DDH diagnosis. There were 192 THAs in 157 patients that did not have adequate demographic data and were excluded from any analysis. The diagnosis was OA in 7792 patients, ON/AVN in 523, RA in 204, and DDH in 60. Those with ON/AVN and RA were then excluded, leaving 9671 THAs in 7852 patients for analysis. Date of birth, age, and gender were gathered from the database of total joint surgeries performed at the authors' institution based on medical records and operation note data at the time of surgery.

The months of the year were divided into winter and non-winter. Winter was defined as October through February, and non-winter as March through September based on local temperatures. The source for the average monthly temperature and precipitation was from the Indiana State Climate Office in Perdue University (https://climate.agry.purdue.edu/climate/facts.asp) (Table 1). Differences by winter and non-winter months were evaluated for both the DDH and OA groups. Temporal variation was analyzed using cosinor analysis (see below).

***Statistical analysis***

Continuous data are reported as the mean ± standard deviation. Categorical data are reported as frequencies or percentages. Statistical differences between continuous variables were analyzed by the student’s *t*-test and between categorical variables by the Pearson’s *χ*2 test. Statistical Analysis System (SAS) version 9.2 (Cary, NC) was used for *t*-tests and categorical analysis where the expected value was set as the null hypothesis. The Statistical Package for Social Sciences (SPSS) version 12.0 for Windows (SPSS Inc., Chicago, IL, USA) was used for further statistical analysis. The data were subjected to cosinor analysis in an effort to mathematically define any seasonal variation when present. Cosinor analysis[8,9] determines the best mathematical fit of the data to an equation defined by F(t) = M + Acos(ωt + φ), where M = the mean level (termed mesor), A = the amplitude of the cosine curve, φ = acrophase (phase angle of the maximum value), ω = the frequency (which for monthly analysis is 360°/12 = 30°), and t = time (which in this case is each month). The overall *P* and r2 value for the distribution are given for the rhythmic pattern described by the cosinor equation for M, A, and φ. The data was analyzed for the entire period of 12 mo as well as decreasing increments of 1 mo. A best monthly fit may not be over a period of 12 mo, but a different time span (*e.g.,* 7 or 6 mo periodicity). Cosinor analyses were performed with ChronoLab 3.0™ software (see Acknowledgement). For all statistical analyses the level of significance was set at *P* < 0.05.

**RESULTS**

The average age at the time of THA was 55.4 ± 13.6 years in the DDH group and 67.9 ± 11.4 years in the OA group. This age difference was statistically significant (*P* < 0.0001). There were 49 females (81.7%) and 11 males (18.3%) in the DDH and 4160 females (53.4%) and 3,632 males (46.6%) in the OA group (*P* < 0.0001). The distribution of birth months for the 7792 OA and 60 DDH patients can be found in Table 1. The number of births in the winter months was much higher than expected in both the DDH [41 of 60 (68.3%) actual, 25 of 60 (41.7%) expected, *P* < 0.0001) and the OA [3371 of 7792 (43.3%) actual, 3247 of 7792 (41.7%) expected, *P* < 0.0052] groups. Cosinor analysis demonstrated an excellent fit for the OA group with a peak on October 1 (Figure 1). There was no good cosinor fit for the DDH group, likely due to the small number.

**DISCUSSION**

This study demonstrates that both adult DDH and OA patients with end stage hip disease are more frequently born in the colder months. This is similar to pediatric DDH in the previous report[3] and adult Japanese OA[7]. Children with DDH undergoing operative treatment demonstrated a peak birth on October 15[3], and is essentially the same as the adult OA patients peak on October 1. Nagamine *et al*[7] noted that adult hip osteoarthritis in the Japanese population was more prevalent when the patient was born in the colder months. The authors subjected their data to cosinor analysis and found a peak on January 20 (Figure 2). The Japanese OA peak of January 20 is identical to the January 26 peak in Japanese children with DDH from Kochi, Japan[3,10].

There are certain limitations to this study. This is a retrospective study and diagnosis of DDH and OA was determined mainly by radiographs and patient history. The diagnosis of OA includes both primary and secondary OA[11]. Patients undergoing THA are usually at the end stage of hip arthritis making it difficult to distinguish primary from secondary OA. Some patients will have shallow acetabulae, but once the joint space disappears at the lateral aspect of the acetabulum, it is impossible to differentiate between primary and secondary OA. The patient’s history may help, but most patients with DDH treated in infancy with a Pavlik harness or other abduction bracing may not be aware of that history; they would not be able to remember it due to their young age at the time of treatment and their parents may not have told them of such treatment. Although the DDH group was small, we still believe that the findings are valid due to the highly significant *P* < 0.0001.

These findings have several possible explanations. First, it is very likely that many of the OA patients had underlying, subtle mild DDH (*e.g.*, minimal acetabular dysplasia). The population of OA, technically the secondary OA, may have just more of DDH than what the authors think. Another possible explanation is that both DDH and OA patients share some common, but as yet unknown, etiology. The possible factors/etiologies that can be invoked are pre-/perinatal maternal factors as well as postnatal environmental factors.

Potential pre-/perinatal factors are a breech birth or difficult delivery[12]. First born and high-birth-weight babies are at an increased risk of DDH, but those do not exhibit seasonal trend[6]. Obstetric pelvic insufficiency shows a seasonal variation with a peak in November to December[13]. In the current study, the authors did not investigate the patients’ birth history (pregnancy number or breech/vertex delivery). Since there been no report showing a relationship between osteoarthritis and breech/difficult delivery, it is most likely that residual DDH (minimal acetabular dysplasia) influenced the incidence of OA.

Another potential perinatal factor is relaxin, which stimulates collagenase and alters connective tissue[14,15]. Collagen metabolism is altered in DDH[16] with increased joint laxity[17]. However, the correlation between DDH and relaxin is controversial[18,19]. In addition, no seasonal variation in relaxin levels has been shown leading to the conclusion that relaxin is not a critical factor. Another possible factor is vitamin D, which is well known to have a seasonal variation with peak levels in the summer[20,21]. Low maternal levels of vitamin D result in low birth weight infants and increased levels in heavy infants[22,23]. A report from Norway suggests hip and knee osteoarthritis is more common in those born in the spring months[24], perhaps due to vitamin D issues.

In conclusion, this study demonstrated that patients undergoing THA for end stage hip disease due to OA or the sequelae of DDH are more commonly born in the colder months. This confirms previous reports noting the same phenomenon in both adult OA and childhood DDH. Further research will be needed to understand this association.

**ACKNOWLEDGEMENTS**

The authors wish to acknowledge the help of Michihiro Kobayashi, MD, PhD, Herman B. Wells Research Center, Department of Pediatrics, Indiana University School of Medicine who extracted the data from the Japanese study by Haginomori[10]. ChronoLab 3.0™ software, designed for use on Macintosh™ computers, cannot be purchased. The software used to perform cosinor analyses was provided through the courtesy of Dr. Artemio Mojón and colleagues, Bioengineering and Chronobiology Labs, ETSI Telecomunicación, University of Vigo, Campus Universitario, Vigo (Pontevedra) 36280, Spain. It can be downloaded from their web site at <http://www.tsc.uvigo.es/BIO/Bioing/References.html>. Please kindly acknowledge their generosity when using this software.

**COMMENTS**

***Background***

Winter birth has been associated with developmental dysplasia of the hip (DDH) and it was reported recently to be related to hip osteoarthritis (OA). This study aims to investigate the epidemiology of DDH and OA according to the registry of total hip arthroplasty in the authors’ institution.

***Research frontiers***

To the authors’ knowledge, this is the first study to present both OA and DDH were associated with winter birth. Reviewing the registry over 40 years, the average age at the surgery was statistically younger in DDH patients than in OA patients. Both DDH and OA patients were more born in winter months than in non-winter months.

***Innovations and breakthroughs***

Several reports have shown DDH is often caused by swaddling during infancy, which was reported to be one reason DDH is associated with winter birth. The study demonstrated OA also was associated with winter birth. The etiology is still unknown why winter birth influences hip OA, but the study will contribute to elucidate the mechanism of OA and the prevention of hip surgery in the future.

***Peer-review***

It is an interesting and well-written article.

**REFERENCES**

1. **Harris WH**. Etiology of osteoarthritis of the hip. *Clin Orthop Relat Res* 1986; **(213)**: 20-33 [PMID: 3780093]
2. **Cooperman DR**, Wallensten R, Stulberg SD. Acetabular dysplasia in the adult. *Clin Orthop Relat Res* 1983; **(175)**: 79-85 [PMID: 6839611 DOI: 10.1097/00003086-198305000-00013]
3. **Loder RT**, Shafer C. Seasonal variation in children with developmental dysplasia of the hip. *J Child Orthop* 2014; **8**: 11-22 [PMID: 24500336 DOI: 10.1007/s11832-014-0558-3]
4. **Anand JK**, Moden I, Myles JW. Incidence of neonatal hip instability: are there seasonal variations? *Acta Orthop Belg* 1992; **58**: 205-208 [PMID: 1632220]
5. **Record RG**, Edwards JH. Environmental influences related to the aetiology of congenital dislocation of the hip. *Br J Prev Soc Med* 1958; **12**: 8-22 [PMID: 13510578 DOI: 10.1136/jech.12.1.8]
6. **Robinson GW**. Birth characteristics of children with congenital dislocation of the hip. *Am J Epidemiol* 1968; **87**: 275-284 [PMID: 5647859]
7. **Nagamine S**, Sonohata M, Kitajima M, Kawano S, Ogawa K, Mawatari M, Hotokebuchi T. Seasonal trends in the incidence of hip osteoarthritis in Japanese patients. *Open Orthop J* 2011; **5**: 134-137 [PMID: 21584203 DOI: 10.2174/1874325001105010134]
8. **Nelson W**, Tong YL, Lee JK, Halberg F. Methods for cosinor-rhythmometry. *Chronobiologia* 1979; **6**: 305-323 [PMID: 548245]
9. **Faure A**, Nemoz C, Claustrat B. A graphical and statistical method for investigation of time series in chronobiology according to the cosinor procedure. *Comput Biol Med* 1990; **20**: 319-329 [PMID: 2257732 DOI: 10.1016/0010-4825(90)90011-D]
10. **Haginomori K**. A statistical study of congenital dislocation of the hip in the south-western part of Kochi prefecture. Shikoku Acta Med 1966; **22**: 112-125
11. **Hoaglund FT**, Steinbach LS. Primary osteoarthritis of the hip: etiology and epidemiology. *J Am Acad Orthop Surg* 2011; **9**: 320-327 [PMID: 11575911 DOI: 10.5435/00124635-200109000-00005]
12. **Fredensorg N**. Observations in children with congenital dislocation of the hip. *Acta Orthop Scand* 1976; **47**: 175-180 [PMID: 944988 DOI: 10.3109/17453677608989714]
13. **Berezin D**. Pelvic insufficiency during pregnancy and after parturition. *Acta Obstet Gynecol Scand Suppl* 1950; **30**: 170-182 [PMID: 14799179 DOI: 10.3109/00016345009164771]
14. **Forst J**, Forst C, Forst R, Heller KD. Pathogenetic relevance of the pregnancy hormone relaxin to inborn hip instability. *Arch Orthop Trauma Surg* 1997; **116**: 209-212 [PMID: 9128773 DOI: 10.1007/BF00393711]
15. **Saugstad LF**. Persistent pelvic pain and pelvic joint instability. *Eur J Obstet Gynecol Reprod Biol* 1991; **41**: 197-201 [PMID: 1936503 DOI: 10.1016/0028-2243(91)90024-F]
16. **Jensen BA**, Reimann I, Fredensborg N. Collagen type III predominance in newborns with congenital dislocation of the hip. *Acta Orthop Scand* 1986; **57**: 362-365 [PMID: 3788503 DOI: 10.3109/17453678608994412]
17. **Carr AJ**, Jefferson RJ, Benson MK. Joint laxity and hip rotation in normal children and in those with congenital dislocation of the hip. *J Bone Joint Surg Br* 1993; **75**: 76-78 [PMID: 8421041 DOI: 10.1097/01241398-199307000-00043]
18. **Hinderaker T**, Daltveit AK, Irgens LM, Udén A, Reikerås O. The impact of intra-uterine factors on neonatal hip instability. An analysis of 1,059,479 children in Norway. *Acta Orthop Scand* 1994; **65**: 239-242 [PMID: 8042471 DOI: 10.3109/17453679408995446]
19. **Vogel I**, Andersson JE, Uldbjerg N. Serum relaxin in the newborn is not a marker of neonatal hip instability. *J Pediatr Orthop* 1998; **18**: 535-537 [PMID: 9661868]
20. **Stryd RP**, Gilbertson TJ, Brunden MN. A seasonal variation study of 25-hydroxyvitamin D3 serum levels in normal humans. *J Clin Endocrinol Metab* 1979; **48**: 771-775 [PMID: 429522 DOI: 10.1210/jcem-48-5-771]
21. **Andersen R**, Mølgaard C, Skovgaard LT, Brot C, Cashman KD, Chabros E, Charzewska J, Flynn A, Jakobsen J, Kärkkäinen M, Kiely M, Lamberg-Allardt C, Moreiras O, Natri AM, O'brien M, Rogalska-Niedzwiedz M, Ovesen L. Teenage girls and elderly women living in northern Europe have low winter vitamin D status. *Eur J Clin Nutr* 2005; **59**: 533-541 [PMID: 15714215 DOI: 10.1038/sj.ejcn.1602108]
22. **Aghajafari F**, Nagulesapillai T, Ronksley PE, Tough SC, O'Beirne M, Rabi DM. Association between maternal serum 25-hydroxyvitamin D level and pregnancy and neonatal outcomes: systematic review and meta-analysis of observational studies. *BMJ* 2013; **346**: f1169 [PMID: 23533188 DOI: 10.1136/bmj.f1169]
23. **Christesen HT**, Elvander C, Lamont RF, Jørgensen JS. The impact of vitamin D in pregnancy on extraskeletal health in children: a systematic review. *Acta Obstet Gynecol Scand* 2012; **91**: 1368-1380 [PMID: 23210535 DOI: 10.1111/aogs.12006]
24. **Fønnebø V**. Arthrosis of the hip and knee: environmental causes in the first year of life? A study of 1405 cases of arthrosis in north Norway 1984-1989. *Arctic Med Res* 1995; **54**: 151-154 [PMID: 7669129]

**P-Reviewer:** Lee SH, Sebestyén A **S-Editor:** Kong JX **L-Editor: E-Editor:**



**Figure 1 The cosinor fit for Indiana patients with hip osteoarthritis and month of birth.** The best fit is represented by the equation: Number of OA births = 646.4 + 55.3(cos(36t -18)-324), where t = 1 is January, 2 is February, 3 is March, etc. This was statistically significant (r2 = 0.56, *P* = 0.025). The peak is October 1 (solid arrow). The data points are the black triangles and the best fit represented by the bold black line. OA: Osteoarthritis.



**Figure 2 The cosinor fit for Japanese patients with osteoarthritis and month of birth.** Data extracted from the study of [7]. The best fit is represented by the equation: number of OA births = 153.7 + 132.7(cos30t-15)-21, where t = 1 is January, 2 is February, 3 is March, etc. This was statistically significant (r2 = 0.83, *P* < 0.001). The peak is January 20 (solid arrow). The data points are the black triangles and the best fit represented by the bold black line. OA: Osteoarthritis.

**Table 1 Month of birth for 7852 patients undergoing total hip arthroplasty for osteoarthritis and sequelae of developmental hip dysplasia**

|  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- |
|  | **DDH** |  | **OA** |  | **Temperature (°F)** | **Precipitation (ins)** |
| Month of birth | *n* | % | *n* | % |  |  |
| January | 10 | 17 | 745 | 9.6 | 26.0 | 2.5 |
| February | 6 | 10 | 622 | 8.0 | 30.5 | 2.3 |
| March | 3 | 5 | 617 | 7.9 | 40.7 | 3.4 |
| April | 3 | 5 | 585 | 7.5 | 51.0 | 3.9 |
| May | 3 | 5 | 582 | 7.5 | 61.4 | 4.5 |
| June | 4 | 7 | 613 | 7.9 | 70.5 | 4.2 |
| July | 3 | 5 | 667 | 8.6 | 74.3 | 4.2 |
| August | 0 | 0 | 675 | 8.7 | 72.2 | 3.9 |
| September | 3 | 5 | 682 | 8.8 | 65.2 | 3.1 |
| October | 14 | 23 | 702 | 9.0 | 53.6 | 3.0 |
| November | 3 | 5 | 610 | 7.8 | 42.3 | 3.4 |
| December | 8 | 13 | 692 | 8.9 | 31.2 | 3.1 |
| Total | 60 | 100 | 7792 | 100 |  | 41.5 |

OA: Osteoarthritis; DDH: Developmental hip dysplasia.