Should paediatricians initiate orthopaedic hip dysplasia referrals for infants with isolated asymmetric skin folds?

C. R. Louer¹
J. D. Bomar²
M. E. Pring²
S. J. Mubarak²
V. V. Upasani³
D. R. Wenger²

Abstract

Purpose Asymmetric skin folds (ASFs) have been linked to developmental dysplasia of the hip (DDH) in select studies, leading to their inclusion in paediatric practice guidelines regarding orthopaedic referral for hip evaluation. The purpose of this study was to investigate the utility of isolated ASFs as a screening tool for DDH in a series of patient referrals evaluated at a single institution.

Methods We performed a retrospective review of consecutive patients between 0 and 12 months of age referred to orthopaedic clinics for isolated ASFs. We recorded radiographic findings (acetabular inclination or alpha angle), diagnosis rendered and treatment administered.

Results A total of 66 patients were included (mean age 6.4 months; 2.47 to 10.76). All patients received pelvic radiographs or ultrasound. In all, 36 patients (55%) were considered normal by their treating physician and 25 (38%) were considered dysplastic and underwent brace treatment. One hip with an isolated ASF was found to have a dislocated hip on radiograph prior to their initial orthopaedic visit. None of the patients in this study have required surgery to date.

Conclusion Using ASFs as a reason for referral led to increased diagnosis of mild dysplasia resulting in orthotic treatment. Thus, in our particular clinical environment, isolated ASFs can be an indicator of mild dysplasia and warrant further workup or referral. Because treatment philosophies regarding recognition and treatment of mild dysplasia vary amongst centres, the value of screening with ASFs likewise depends on the treating orthopaedic surgeon’s threshold for treatment of mild dysplasia.

Level of evidence: Level IV- Retrospective

Cite this article: Louer CR, Bomar JD, Pring ME, Mubarak SJ, Upasani VV, Wenger DR. Should paediatricians initiate orthopaedic hip dysplasia referrals for infants with isolated asymmetric skin folds? J Child Orthop 2019;13:593-599. DOI: 10.1302/1863-2548.13.190090

Keywords: DDH referral; hip dysplasia; asymmetric thigh folds

Introduction

Asymmetric skin folds (ASFs) around the hips in children are often considered an early clinical sign for diagnosing developmental dysplasia of the hip (DDH). Early reports assert a relationship between subluxated or dislocated hips with additional thigh or gluteal folds.¹ Subsequent studies have expanded this association to reduced hips with additional thigh or gluteal folds.¹ Several studies claim that ASFs are an indicator of DDH due to their increased prevalence in patients with DDH compared with the normal population (27% to 83%²,³,⁵ rate in DDH patients versus 20%⁵⁻⁷). However, the significant variation in these rates calls into question the definition of asymmetry and the degree needed to be declared abnormal. Furthermore, these studies fail to include a control group that is not pre-screened that would allow a direct comparison of ASF rates using identical criteria. Adding to the confusion are similar studies with comparable flaws that contrarily do not demonstrate a large difference between the rates of ASFs in normal versus DDH populations.⁶ Presently, the question as to the utility of ASFs as a diagnostic indicator of DDH remains unresolved.

Despite the absence of conclusive evidence for an association of ASFs with DDH, guidelines for general paediatricians, who perform most of the early DDH screenings in the United States, continue to instruct that ‘asymmetric thigh or buttock creases’ are a physical exam finding that may indicate DDH.⁸,⁹ In recent years, our orthopaedic surgery group has anecdotally noticed an increased referral rate of infants with all types of ASFs. Given the

¹ University of North Carolina, Chapel Hill, North Carolina, USA
² Rady Children’s Hospital, San Diego, California, USA

Correspondence should be sent to V. Upasani, 3020 Children’s Way, MC5062, San Diego, CA 92123, USA.
E-mail: vupasanirchsdorg
high prevalence of ASFs within the normal population, we questioned the utility of referring patients for DDH evaluation on the basis of ASFs alone.

The purpose of this study, therefore, was to investigate our institution’s recent experience with infants referred for DDH assessment due to isolated ASFs. Rady Children’s Hospital, San Diego is a tertiary referral centre at an academically affiliated stand-alone children’s hospital with more than 1500 patients seen for DDH annually. We reviewed all patients referred for DDH evaluation because of an isolated finding of asymmetric gluteal, inguinal or thigh folds, and excluded those who were referred for any other reason. We catalogued the diagnosis and treatment of these patients to understand whether isolated ASFs led to the diagnosis of DDH within our practice environment. We hypothesized that infants referred for isolated ASFs would have typical hip development and would rarely have a subluxated or dislocated hip.

Materials and methods
The institutional review board approved this study. The electronic medical record was screened for patients with a primary referral reason of ‘Asymmetric Thigh Folds’ as entered by the orthopaedic practitioner at the time of the visit. Search dates were from May 2014 (when we began cataloguing referral reason) to February 2017.

The most comprehensive literature on ASFs has described asymmetric inguinal folds as being associated with an increased incidence of hip subluxation or dislocation in infants.5 The association between middle or distal thigh ASFs with hip dysplasia is less clear.10 Because the referring physicians and clinics do not seem to distinguish between the particular locations of ASFs, we have included all varieties in this study. We use the term asymmetric skin folds (or acronym ASFs) to describe any asymmetry of surface anatomy around the hip joint.

A retrospective chart review of all patients was conducted and the referral reason was verified. Patients were excluded if older than one year, or if there was a referral reason or diagnosis in addition to ASFs. We recorded demographic information, imaging type, diagnosis rendered (normal, dysplasia, subluxation/dislocation) and subsequent treatment. Anteroposterior and frog view pelvis radiographs were graded for the acetabular inclination (AI) by a single grader. The highest AI among either hip on either radiographic view was used for analysis. Subjects that were evaluated with an ultrasound at initial visit were graded using alpha angle. Alpha angle was measured on the coronal view for each hip and the lowest alpha angle among the two hips was used for analysis. Based on the work of Tönnis11 and Graf12 AI and alpha angle were used to grade the acetabulum as normal, mild DDH or DDH (Table 1). Descriptive statistics were performed for all outcome measures using Microsoft Excel (version 14; Redmond, Washington), except for distribution testing and analysis of variance testing between groups for AI which was evaluated using SPSS (version 24; IBM, Armonk, New York).

Results
A total of 66 patients were included and were seen by any one of five treating surgeons, all fellowship-trained in paediatric orthopaedics with experience in treating DDH. Mean age at initial presentation was 6.4 months (SD 2.1; 2.47 to 10.76). Mean AI was 29.5° (SD 4.2°; 21° to 45°). Four subjects were initially evaluated with an ultrasound exam, their ages ranging from 2.47 months to 2.99 months and their alpha angles ranging from 55° to 65°. In all, 83% (55/66) of our cohort were female. A full list of our cohort characteristics can be found in Table 2.

A total of 30 patients (45%) were diagnosed with acetabular dysplasia by their physician at time of initial evaluation.DDH, developmental dysplasia of the hip

| Image type (measurement) | Sex     | Age, mths | Side | Normal | Mild DDH | DDH |
|--------------------------|---------|-----------|------|--------|----------|-----|
| Radiograph (acetabular inclination) | Female  | 3 & 4     | R    | < 31°  | 31° to 36° | > 36° |
|                          |         |           | L    | < 33°  | 33° to 39° | > 39° |
|                          |         | 5 & 6     | R    | < 27°  | 27° to 32° | > 32° |
|                          |         |           | L    | < 29°  | 29° to 34° | > 34° |
|                          |         | 7 to 12   | R    | < 25°  | 25° to 29° | > 29° |
|                          |         |           | L    | < 27°  | 27° to 31° | > 31° |
|                          | Male    | 3 & 4     | R    | < 28°  | 28° to 32° | > 32° |
|                          |         |           | L    | < 29°  | 29° to 34° | > 34° |
|                          |         | 5 & 6     | R    | < 24°  | 24° to 29° | > 29° |
|                          |         |           | L    | < 27°  | 27° to 32° | > 32° |
|                          |         | 7 to 9    | R    | < 25°  | 25° to 29° | > 29° |
|                          |         |           | L    | < 25°  | 25° to 30° | > 30° |
|                          |         | 10 to 12  | R    | < 23°  | 23° to 27° | > 27° |
|                          |         |           | L    | < 25°  | 25° to 29° | > 29° |
| Ultrasound (alpha angle) | M or F  | 0 to 3    | R or L | > 60° | 50° to 59° | < 49° |

DDH, developmental dysplasia of the hip

Table 1 Radiographic classification
| Patient | Age, mths | Sex | Exam | Shenton's Line | Acetabular corner | X-ray prior to orthopaedist visit | AI | Alpha angle | Dx based on x-ray/US | Dx rendered at visit | Treatment |
|---------|-----------|-----|------|---------------|------------------|-----------------------------------|----|-------------|---------------------|---------------------|-----------|
| 1       | 2         | F   | US   | NA            | Blunt            | No prior x-ray                    | NA | 55°         | Mild DDH             | Normal              | Not treated |
| 2       | 3         | F   | US   | NA            | Sharp            | No prior x-ray                    | NA | 60°         | WNL                 | Normal              | Not treated |
| 3       | 3         | F   | X-ray| Unbroken     | Sharp            | No prior x-ray                    | 26 | NA          | WNL                 | Normal              | Not treated |
| 4       | 3         | F   | X-ray| Unbroken     | Blunt            | No prior x-ray                    | 31 | NA          | Mild DDH             | Acetabular dysplasia| Treated   |
| 5       | 3         | F   | X-ray| Unbroken     | Sharp            | No prior x-ray                    | 30 | NA          | WNL                 | Normal              | Not treated |
| 6       | 3         | F   | X-ray| Unbroken     | Blunt            | No prior x-ray                    | 38 | NA          | Mild DDH             | Normal              | Not treated |
| 7       | 4         | F   | X-ray| Unbroken     | Blunt            | Prior x-ray                       | 27 | NA          | WNL                 | Normal              | Not treated |
| 8       | 4         | F   | X-ray| Unbroken     | Blunt            | No prior x-ray                    | 32 | NA          | Mild DDH             | Normal              | Not treated |
| 9       | 4         | F   | X-ray| Unbroken     | Sharp            | No prior x-ray                    | 31 | NA          | Mild DDH             | Normal              | Not treated |
| 10      | 5         | F   | X-ray| Unbroken     | Sharp            | No prior x-ray                    | 29 | NA          | Mild DDH             | Normal              | Not treated |
| 11      | 5         | F   | X-ray| Unbroken     | Blunt            | No prior x-ray                    | 29 | NA          | Mild DDH             | Acetabular dysplasia| Not treated |
| 12      | 5         | F   | X-ray| Unbroken     | Sharp            | No prior x-ray                    | 25 | NA          | WNL                 | Acetabular dysplasia| Treated   |
| 13      | 5         | F   | X-ray| Unbroken     | Sharp            | No prior x-ray                    | 27 | NA          | Mild DDH             | Normal              | Not treated |
| 14      | 5         | F   | X-ray| Unbroken     | Sharp            | Prior x-ray                       | 32 | NA          | Mild DDH             | Normal              | Not treated |
| 15      | 5         | F   | X-ray| Unbroken     | Blunt            | Prior x-ray                       | 32 | NA          | Mild DDH             | Acetabular dysplasia| Treated   |
| 16      | 5         | F   | X-ray| Unbroken     | Sharp            | Prior x-ray                       | 31 | NA          | Mild DDH             | Acetabular dysplasia| Treated   |
| 17      | 5         | F   | X-ray| Unbroken     | Sharp            | No prior x-ray                    | 24 | NA          | WNL                 | Normal              | Not treated |
| 18      | 5         | F   | X-ray| Unbroken     | Sharp            | Prior x-ray                       | 36 | NA          | DDH                 | Acetabular dysplasia| Treated   |
| 19      | 5         | F   | X-ray| Unbroken     | Sharp            | No prior x-ray                    | 30 | NA          | Mild DDH             | Normal              | Not treated |
| 20      | 6         | F   | X-ray| Unbroken     | Sharp            | No prior x-ray                    | 36 | NA          | DDH                 | Acetabular dysplasia| Treated   |
| 21      | 6         | F   | X-ray| Unbroken     | Blunt            | Prior x-ray                       | 30 | NA          | Mild DDH             | Acetabular dysplasia| Not treated |
| 22      | 6         | F   | X-ray| Unbroken     | Blunt            | Prior x-ray                       | 22 | NA          | WNL                 | Normal              | Not treated |
| 23      | 6         | F   | X-ray| Unbroken     | Sharp            | Prior x-ray                       | 30 | NA          | Mild DDH             | Normal              | Not treated |
| 24      | 6         | F   | X-ray| Unbroken     | Sharp            | Prior x-ray                       | 31 | NA          | Mild DDH             | Normal              | Not treated |
| 25      | 6         | F   | X-ray| Unbroken     | Blunt            | Prior x-ray                       | 32 | NA          | Mild DDH             | Normal              | Not treated |
| 26      | 6         | F   | X-ray| Unbroken     | Blunt            | Prior x-ray                       | 36 | NA          | DDH                 | Acetabular dysplasia| Treated   |
| 27      | 6         | F   | X-ray| Unbroken     | Blunt            | Prior x-ray                       | 31 | NA          | Mild DDH             | Acetabular dysplasia| Treated   |
| 28      | 6         | F   | X-ray| Broken       | Sharp            | No prior x-ray                    | 29 | NA          | Mild DDH             | Acetabular dysplasia| Treated   |
| 29      | 6         | F   | X-ray| Broken       | Sharp            | No prior x-ray                    | 30 | NA          | Mild DDH             | Acetabular dysplasia| Treated   |
| 30      | 7         | F   | X-ray| Unbroken     | Blunt            | Prior x-ray                       | 31 | NA          | DDH                 | Acetabular dysplasia| Treated   |
| 31      | 7         | F   | X-ray| Unbroken     | Sharp            | No prior x-ray                    | 21 | NA          | WNL                 | Normal              | Not treated |
| 32      | 7         | F   | X-ray| Unbroken     | Sharp            | No prior x-ray                    | 22 | NA          | WNL                 | Normal              | Not treated |
| 33      | 7         | F   | X-ray| Broken       | Blunt            | Prior x-ray                       | 34 | NA          | DDH                 | Acetabular dysplasia| Treated   |
| 34      | 7         | F   | X-ray| Unbroken     | Sharp            | Prior x-ray                       | 24 | NA          | WNL                 | Normal              | Not treated |
| 35      | 7         | F   | X-ray| Unbroken     | Sharp            | No prior x-ray                    | 28 | NA          | Mild DDH             | Normal              | Not treated |
| 36      | 7         | F   | X-ray| Broken       | Blunt            | Prior x-ray                       | 45 | NA          | DDH                 | Dislocation         | Treated   |
| 37      | 7         | F   | X-ray| Unbroken     | Blunt            | No prior x-ray                    | 21 | NA          | WNL                 | Normal              | Not treated |
| 38      | 7         | F   | X-ray| Unbroken     | Sharp            | Prior x-ray                       | 30 | NA          | DDH                 | Normal              | Not treated |
| 39      | 7         | F   | X-ray| Unbroken     | Sharp            | Prior x-ray                       | 32 | NA          | DDH                 | Acetabular dysplasia| Treated   |
| 40      | 7         | F   | X-ray| Unbroken     | Blunt            | Prior x-ray                       | 30 | NA          | DDH                 | Normal              | Not treated |
| 41      | 8         | F   | X-ray| Unbroken     | Sharp            | No prior x-ray                    | 26 | NA          | Mild DDH             | Acetabular dysplasia| Treated   |
| 42      | 8         | F   | X-ray| Unbroken     | Blunt            | Prior x-ray                       | 30 | NA          | Mild DDH             | Acetabular dysplasia| Treated   |
| 43      | 8         | F   | X-ray| Broken       | Blunt            | No prior x-ray                    | 33 | NA          | DDH                 | Acetabular dysplasia| Treated   |
| 44      | 8         | F   | X-ray| Unbroken     | Sharp            | Prior x-ray                       | 33 | NA          | DDH                 | Normal              | Not treated |
| 45      | 8         | F   | X-ray| Unbroken     | Blunt            | Prior x-ray                       | 33 | NA          | DDH                 | Acetabular dysplasia| Treated   |
| 46      | 8         | F   | X-ray| Unbroken     | Blunt            | Prior x-ray                       | 28 | NA          | Mild DDH             | Acetabular dysplasia| Treated   |
| 47      | 8         | F   | X-ray| Broken       | Blunt            | Prior x-ray                       | 29 | NA          | Mild DDH             | Acetabular dysplasia| Treated   |
| 48      | 8         | F   | X-ray| Unbroken     | Blunt            | No prior x-ray                    | 27 | NA          | Mild DDH             | Normal              | Not treated |
| 49      | 9         | F   | X-ray| Unbroken     | Blunt            | No prior x-ray                    | 30 | NA          | Mild DDH             | Acetabular dysplasia| Treated   |
### Table

| Patient | Age, mths | Sex | Exam | Shenton’s Line | Acetabular corner | X-ray prior to orthopaedist visit | Al | Alpha angle | Dx based on x-ray/US | Dx rendered at visit | Treatment |
|---------|-----------|-----|------|---------------|------------------|-------------------------------|----|-------------|---------------------|----------------------|-----------|
| 50      | 10        | F   | X-ray| Unbroken      | Sharp            | Prior x-ray                  | 25 | NA          | Mild DDH            | Normal               | Not treated |
| 51      | 10        | F   | X-ray| Unbroken      | Sharp            | Prior x-ray                  | 29 | NA          | Mild DDH            | Acetabular dysplasia| Not treated |
| 52      | 10        | F   | X-ray| Unbroken      | Blunt            | Prior x-ray                  | 29 | NA          | Mild DDH            | Acetabular dysplasia| Treated    |
| 53      | 10        | F   | X-ray| Unbroken      | Blunt            | Prior x-ray                  | 26 | NA          | Mild DDH            | Normal               | Not treated |
| 54      | 10        | F   | X-ray| Unbroken      | Sharp            | No prior x-ray               | 25 | NA          | WNL                 | Normal               | Not treated |
| 55      | 11        | F   | X-ray| Unbroken      | Blunt            | No prior x-ray               | 28 | NA          | Mild DDH            | Normal               | Not treated |
| 56      | 3         | M   | US   | NA            | Blunt            | No prior x-ray               | 58 | 65°         | WNL                 | Normal               | Not treated |
| 57      | 3         | M   | US   | NA            | Sharp            | No prior x-ray               | 33 | NA          | DDH                 | Acetabular dysplasia| Treated    |
| 58      | 5         | M   | X-ray| Unbroken      | Blunt            | No prior x-ray               | 27 | NA          | Mild DDH            | Acetabular dysplasia| Treated    |
| 59      | 6         | M   | X-ray| Unbroken      | Blunt            | Prior x-ray                  | 35 | NA          | DDH                 | Normal               | Not treated |
| 60      | 6         | M   | X-ray| Unbroken      | Sharp            | Prior x-ray                  | 27 | NA          | Mild DDH            | Acetabular dysplasia| Treated    |
| 61      | 7         | M   | X-ray| Broken        | Sharp            | No prior x-ray               | 30 | NA          | Mild DDH            | Acetabular dysplasia| Not treated |
| 62      | 7         | M   | X-ray| Unbroken      | Blunt            | Prior x-ray                  | 27 | NA          | Mild DDH            | Acetabular dysplasia| Not treated |
| 63      | 7         | M   | X-ray| Unbroken      | Blunt            | No prior x-ray               | 30 | NA          | Mild DDH            | Normal               | Not treated |
| 64      | 8         | M   | X-ray| Unbroken      | Blunt            | No prior x-ray               | 24 | NA          | WNL                 | Normal               | Not treated |
| 65      | 8         | M   | X-ray| Unbroken      | Sharp            | No prior x-ray               | 28 | NA          | Mild DDH            | Acetabular dysplasia| Treated    |
| 66      | 10        | M   | X-ray| Unbroken      | Blunt            | Prior x-ray                  | 28 | NA          | Mild DDH            | Acetabular dysplasia| Treated    |

*rounded to the nearest month
**treatment initiated at second visit
AI, acetabular inclination; Dx, diagnosis; US, ultrasound; NA, not available; DDH, developmental dysplasia of the hip; WNL, within normal limits

### Discussion

Our study sought to investigate the utility of using ASFs as criteria for DDH referral by paediatricians. We identified and analyzed a cohort of patients that were referred solely due to ASFs who conceivably would not have been evaluated by an orthopaedist without this finding or if the policy of advising referral on the basis of ASFs did not exist. One of these patients had a dislocated hip. Interestingly, although 79% of our cohort was found to have acetabular dysplasia based on strict radiographic classification, only 38% of our cohort had dysplasia significant enough to warrant treatment in the eyes on their treating physician (Fig. 1).

DDH in the newborn remains one of the few orthopaedic conditions in the United States for which routine screening is recommended. The concept is that a poor outcome (due to a late-recognized hip dislocation) can be prevented by early identification and intervention, as early treatments are less risky to the developing hip. Screening methods are not without controversy, however, as there is question as to their efficacy in reducing the rate of missed dislocations and there is evidence that screening significantly increases treatment rates. In addition, there is no cost-effective benchmark test for DDH, which turns DDH screening into an exercise in risk-stratification for the paediatrician. The ideal screening test would have a high sensitivity to limit the number of false negative tests (missed diagnoses), as well as high specificity to limit the number of false positive tests (unnecessary referrals to orthopaedic surgeons).

The utility of ASFs has been asserted and refuted throughout the DDH literature. Presumably, this is due to the varying definition of what constitutes an abnormal...
fold and the practitioner’s sensitivity to this finding. Most estimates indicate that even normal infants have ASFs at about a 20% rate. The most rigorous study of ASFs comes from Ando and Gotoh from Japan, who studied abnormal inguinal creases in the frog-leg position of 2111 infants referred to orthopaedics and correlated their findings to the radiographic diagnosis. They found an ASF rate of 23.8%, and found that ASFs were present in all 12 patients with dislocations or subluxations. Acetabular dysplasia without subluxation was diagnosed in 17 patients, yet only 41.7% of these patients exhibited ASFs. Their study demonstrated that ASFs are a sensitive test for dislocation or subluxation, but lack sensitivity in detecting acetabular dysplasia. Furthermore, with a 23.8% rate of ASFs in this large cohort, specificity was exceptionally poor as there were 470 patients with ASFs who did not have the disease. Using a prospectively collected sample of infants referred for DDH evaluation in the United Kingdom, Anderton et al studied a 105-patient subgroup with ASFs to investigate the diagnostic value of the ASFs. Only two (2%) of these patients were found to have pathological DDH (subluxation or dislocation), and both cases had other physical exam findings present (limited abduction and Galeazzi sign) to diagnose a dislocation. The authors conclude that ASFs do not add value to the DDH examination, as these other exam findings are just as sensitive, with more specificity.

In the present study, we took a different approach than the aforementioned studies by investigating the fate of those patients who are referred solely because of ASF findings. This theoretically allows us to evaluate the additive benefit of ASF screening by eliminating contributions of other clinical factors. Like the Anderton et al study, we did not associate hip subluxation or dislocation with ASFs in isolation. Distinct from prior reports, however, we found that 38% of patients referred with ASFs were treated for acetabular dysplasia with hip orthoses. The reason for this discrepancy in treatment rates is likely due to variability in the threshold to diagnose and treat dysplasia. In our study, those that were treated for dysplasia had an average acetabular index of 29.5°, which is higher than expected at six months of age (girls ~24°, boys ~22°). With growing recognition of the prevalence and morbidity of dysplasia in young adult hips, there are many paediatric orthopaedists who favour early treatment of mild dysplasia, both by orthoses and surgery, when necessary. Undoubtedly, there are patients within our study cohort who would have normalized without treatment, yet with non-invasive and low-risk treatment options such as orthoses, it is not surprising that many surgeons opt to treat mild dysplasia to ensure its resolution.

This retrospective and small study does have noteworthy limitations based on its design and the nature of the
subject. The approach of investigating patients who are referred to orthopaedists specifically because of ASFs, while offering a critical look at those patients, does limit the number of patients available for evaluation. Finding only 66 patients referred for isolated ASFs out of such a large potential pool demonstrates that our initial over-referral concerns were likely unfounded. In all, 44% of our cohort had a radiograph prior to their orthopaedic visit. We cannot determine if those radiographs were reviewed prior to the paediatrician referring the patient to orthopaedics, or if they were simply ordered to be performed prior to the orthopaedic visit because the paediatrician understood that radiographs would be needed. If some paediatricians routinely wait for a radiologist’s reading prior to referral to orthopaedics, we have no way of knowing how many radiographs were ordered by paediatricians because of ASFs, but were not referred to orthopaedists because the radiologist’s reading was negative. This may inflate the percentage of subjects with ASFs that require treatment. However, our orthopaedic surgeons were found to disagree with the radiology readings 42% of the time which may offset the risk of sample bias. Another limitation is the short follow-up interval. While the goal of this study was to document the initial treatment of screened patients with isolated ASF, there is no doubt that long-term follow-up of these patients could allow us to see the effects of our treatment decisions. In fact, a major limitation to all studies on this topic is the poor understanding of the natural history of mild dysplasia. Until the orthopaedic community gains the ability to differentiate between those patients with mild dysplasia who warrant treatment and the great majority that will improve on their own, this uncertainty will remain pervasive in this and other studies.

Clearly the North American medical-legal climate makes definitive diagnosis and treatment of infantile DDH problematic for multiple medical specialists including paediatricians, radiologists and orthopaedic surgeons. Paediatricians, averse to missing a DDH diagnosis and diligently following their academy guidelines, search carefully for findings such as ASFs and then refer for imaging studies. The radiologist, fearing legal consequences, is more likely to err on the side of over-diagnosis when encountering a borderline case. Finally, the orthopaedic surgeon, faced with anxious caretakers and a diagnosis of dysplasia already in the printed record, may be more inclined to treat the mild dysplasia.

How our experience should be applied is likely related to the economic and cultural climate in which one practices. In circumstances that necessitate an economical approach to diagnosis and treatment, where the concern is only detection of subluxation/dislocation, ASFs alone seem to offer no additional value. If the practice environment warrants that the physicians recognize (and potentially treat) all dysplasia, including mild cases, then isolated ASFs can identify patients who would otherwise escape detection. If this philosophy is to be utilized, the literature would suggest that focusing on more proximal or inguinal asymmetry offers greater diagnostic value.

After careful analysis of our results and consideration of limitations, we conclude that there can be utility in the use of ASFs as a screening criteria for acetabular dysplasia, depending on the orthopaedist’s threshold for treatment. Referring paediatricians should continue to use this as only a single criterion among many, recognizing that ASFs in infancy are a very common finding with limited specificity. Orthopaedic surgeons might consider their threshold for dysplasia treatment when advising their referral base on the utility of ASFs in their particular practice.

Future studies on this subject would require prospective evaluation of all infants to characterize their specific skin fold anomaly so that we may better understand if particular patterns enhance diagnostic ability. Also, further prospective evidence regarding late outcomes of mild dysplasia with and without treatment would greatly improve our understanding of DDH screening practices.

Received 10 June 2019; accepted after revision 18 September 2019.

COMPLIANCE WITH ETHICAL STANDARDS

FUNDING STATEMENT
No benefits in any form have been received or will be received from a commercial party related directly or indirectly to the subject of this article.

OA LICENCE TEXT
This article is distributed under the terms of the Creative Commons Attribution-Non Commercial 4.0 International (CC BY-NC 4.0) licence (https://creativecommons.org/licenses/by-nc/4.0/) which permits non-commercial use, reproduction and distribution of the work without further permission provided the original work is attributed.

ETHICAL STATEMENT
Ethical approval: All procedures performed in studies involving human participants were in accordance with the ethical standards of the institutional and/or national research committee and with the 1964 Helsinki declaration and its later amendments or comparable ethical standards.

Informed consent: This retrospective study was approved by our governing institutional review board and awarded a waiver of consent; therefore consent was not obtained by subjects whose medical information was used in this study.

ICMJE CONFLICT OF INTEREST STATEMENT
SJM reports paid consultancy for Rhino Pediatric Orthopedic Designs, Inc., outside the submitted work.

VVU reports work as a paid presenter or speaker for BroadWater, DePuy, A Johnson & Johnson Company, Nuvasive and OrthoPediatrics; research support for EOS Imaging; paid consultancy for Globus Medical and OrthoPediatrics; stock or stock options in Imagen; research support for Pacira; publishing royalties, financial or material support from Wolters Kluwer Health – Lippincott Williams & Wilkins, outside the submitted work.
DDH AND ASYMMETRIC SKIN FOLDS

DRW reports paid consultancy for Rhino Pediatric Orthopedic Designs, Inc., outside the submitted work; publishing royalties, financial or material support from Wolters Kluwer Health – Lippincott Williams & Wilkins, outside the submitted work. All other authors declare no conflict of interest.

AUTHOR CONTRIBUTIONS
CRL: Study design, Data collection, Data interpretation, Manuscript preparation.
JDB: Study design, Data analysis, Data interpretation, Manuscript preparation.
MEP: Study design, Manuscript preparation.
SJM: Study design, Manuscript preparation, Data interpretation.
VVU: Study design, Manuscript preparation, Data interpretation.
DRW: Study design, Manuscript preparation, Data interpretation.

REFERENCES
1. Dimon JH III. Observation of the gluteal fold in hip dysplasia. Clin Orthop Relat Res 1974;103:19.
2. Abu Hassan FO, Shannak A. Associated risk factors in children who had late presentation of developmental dysplasia of the hip. J Child Orthop 2007;1:205-210.
3. Köse N, Omeroğlu H, Ozyurt B, et al. Our three-year experience with an ultrasonographic hip screening program conducted in infants at 3 to 4 weeks of age. Acta Orthop Traumatol Turc 2006;40:285-290.
4. Omeroğlu H, Koparal S. The role of clinical examination and risk factors in the diagnosis of developmental dysplasia of the hip: a prospective study in 188 referred young infants. Arch Orthop Trauma Surg 2001;121:7-11.
5. Ando M, Gotoh E. Significance of inguinal folds for diagnosis of congenital dislocation of the hip in infants aged three to four months. J Pediatr Orthop 1990;10:331-334.
6. Stein-Zamir C, Volovik I, Rishpon S, Sabri R. Developmental dysplasia of the hip: risk markers, clinical screening and outcome. Pediatr Int 2008;50:341-345.
7. Stoffelen D, Urlus M, Molenaers G, Fabry G. Ultrasound, radiographs, and clinical symptoms in developmental dislocation of the hip: a study of 170 patients. J Pediatr Orthop B 1995;4:194-199.
8. American Academy of Pediatrics. Clinical practice guideline: early detection of developmental dysplasia of the hip. Committee on Quality Improvement, Subcommittee on Developmental Dysplasia of the Hip. Pediatrics 2000;105:896-905.
9. Shaw BA, Segal LS, Section on Orthopedics. Evaluation and referral for developmental dysplasia of the hip in infants. Pediatrics 2016;138:e20163107.
10. Barlow TG. Early diagnosis and treatment of congenital dislocation of the hip. J Bone Joint Surg [Br] 1962;44-B:292-301.
11. Tönnis D. Normal values of the hip joint for the evaluation of X-rays in children and adults. Clin Orthop Relat Res 1976;119:39-47.
12. Graf R. Hip sonography: diagnosis and management of infant hip dysplasia. 2nd ed. Berlin, Heidelberg: Springer-Verlag, 2006.
13. Shorter D, Hong T, Osborn DA. Cochrane review: screening programmes for developmental dysplasia of the hip in newborn infants. Evid Based Child Health 2013;8:11-54.
14. Price KR, Dove R, Hunter JB. Current screening recommendations for developmental dysplasia of the hip may lead to an increase in open reduction. Bone Joint J 2013;95-B:846-850.
15. Anderton MJ, Hastie GR, Paton RW. The positive predictive value of asymmetrical skin creases in the diagnosis of pathological developmental dysplasia of the hip. Bone Joint J 2018;100-B:675-679.
16. Scoles PV, Boyd A, Jones PK. Roentgenographic parameters of the normal infant hip. J Pediatr Orthop 1987;7:656-663.
17. Wenger DR. Is there a role for acetabular dysplasia correction in an asymptomatic patient? J Pediatr Orthop 2013;33(suppl 1):S8-S12.
18. Gans I, Flynn JM, Sankar WN. Abduction bracing for residual acetabular dysplasia in infantile DDH. J Pediatr Orthop 2013;33:714-718.
19. Roposch A, Liu LQ, Hefti F, Clarke NMP, Wedge JH. Standardized diagnostic criteria for developmental dysplasia of the hip in early infancy. Clin Orthop Relat Res 2011;469:3451-3461.
20. Bialik V, Bialik GM, Blazer S, et al. Developmental dysplasia of the hip: a new approach to incidence. Pediatrics 1999;103:93-99.
21. Mladenov K, Dora C, Wicart P, Seringe R. Natural history of hips with borderline acetabular index and acetabular dysplasia in infants. J Pediatr Orthop 2002;22:607-612.