Failure by Immune Reaction to Metal Debris in Total Joint Arthroplasty of Hip and Knee

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Abstract

Background: We propose a clinical classification for Failure by Immune Reaction to Metal debris (FIRM) in Total Joint Arthroplasty (TJA) of hip and knee. We also developed an evidence based diagnostic scoring system and estimated the treatment threshold for FIRM.

Methods: Pub Med and Embase search engines were used to identify original articles. We classified FIRM and identified the individual diagnostic criteria for each type of FIRM. For each individual diagnostic criterion we estimated the pooled Diagnostic Odds Ratio (DOR+) and the individual discriminatory FIRM (id FIRM) scores. From these scores total discriminatory FIRM (TdFIRM) scores were calculated.

Results: We identified a total of 39 original articles for meta-analysis. Based on predominant symptom of clinical presentation, we classified FIRM in to two types i.e., FIRMtype1 and FIRMtype2. We identified 8 individual diagnostic criteria for FIRMtype1, and FIRMtype2 and for each type we estimated the pooled diagnostic odds ratio (DOR+) and the individual discriminatory FIRM (id FIRM) scores. TdFIRM score scores for FIRMtype1 and FIRMtype2 were 4.83 and 4.85 respectively. Treatment threshold for FIRMtype1 and FIRMtype2 were estimated to be 3.38 and 3.39 respectively.

Conclusion: This meta-analysis provides a clinically useful tool for decision making when evaluating a patient suspected to have developed clinical complication by immune reaction to metal debris from arthroplasty. Future studies on FIRM should utilize this scoring system in decision making and critically evaluate its validity.

Keywords: Total joint arthroplasty; Osteolysis; Failure by immune reaction to metal debris; Metal allergy; Metal hypersensitivity

Introduction

The terms metal allergy and metal hypersensitivity are used interchangeably in the current literature to describe a spectrum of complications caused by immune reaction to metal debris generated in total joint arthroplasty. At one end of the spectrum is dermatitis. The dermatitis caused by immune reaction to metal debris may be new in onset or exacerbation of previous lesions and, can be localized or generalized manifestation.

At the other end of the spectrum is a consequence of local immune reaction to metal debris generated by total joint arthroplasty presenting clinically as a variable combination of pain in the joint, cystic lesions around the joint, aseptic loosening of implants and instability of the joint. Various terminologies exist to describe histopathological and imaging findings in patients with pain in the joint suspected to be from local reaction to metal debris after total joint arthroplasty. Willert et al termed the histopathological findings in these patients as Aseptic Lymphocyte-dominated Vasculitis-Associated Lesion (ALVAL) and LymphocYte-Dominated Immunological Answer (LYDIA) [1]. Another histopathological term ‘Metallosis’ was used by Korovesis et al to describe similar findings [2]. Pandit et al used the term ‘Pseudotumour’ to describe cystic and solid masses associated with resurfacing devices [3]. Longton et al described an umbrella term Adverse Reaction to Metal Debris (ARMRD) to describe clinical failures of hip joint arthroplasty presenting with pain, a large sterile effusion and/or macroscopic necrosis [4].

For the purpose of this study, we used the acronym ‘FIRM’ (Failure by Immune Reaction to Metal debris) to describe all the clinical failures of Total Joint Arthroplasty (TJA) in hip and also knee joint due to immune reaction to metal debris. The objectives of this study are twofold. One is to develop a classification system for FIRM in TJA of hip and knee that is based on distinctive clinical manifestations. Second is to develop a diagnostic scoring system to facilitate the clinician to make an accurate, evidence based diagnosis.

Materials and Methods

In January 2015, a Pub Med and Embase search was performed independently using two search terms ‘Metal Allergy Joint Replacement’ (method 1) and ‘Metal Hypersensitivity Joint Replacement’ (method 2) for record identification. The search term that yielded the maximum number of results among the two was used for records identification from that particular search engine. The duplicates among Pub Med and Embase search were identified and removed. The selected records were screened by reviewing...
the abstract for clinical data on TJA patients with FIRM and were included for full-text review. If abstract is not available for a record, it was selected for full-text review by default. Of all the records selected for full-text review, final analysis included only the articles with clinical data on TJA patients with FIRM who had a negative work-up for the possibility of infection or neoplasm (primary or metastatic) and could successfully be managed by removal of implants and reinsertion of new implants that generate no metal debris or by pseudoarthrosis (positive control). In addition, we looked carefully to identify such additional records which we may have missed during Pub Med or Embase search but have the data that we are interested in by reviewing the references section of all the full-text reviewed records and if such record is found, was included in the systematic review and meta analysis.

Part I: FIRM clinical classification

By systematic review of the articles selected for final analysis, we identified all the described complications due to FIRM in patients with TJA and grouped them depending on the most commonly observed patterns of clinical presentation.

Part II: FIRM diagnostic scoring system

A meta-analysis was performed to develop FIRM scoring system using the data from the full-text reviewed records. The first step was calculation of pooled diagnostic odds ratio (also called likelihood ratio) of all the individual diagnostic criteria for each defined type of FIRM using the formula,

\[
DOR^+ = \frac{\text{Most commonly observed result of a diagnostic criteria} + 0.5}{\text{Least commonly observed result of a diagnostic criteria} + 0.5}
\]

Addition of 0.5 to all counts is a validated statistical method to obtain a definable value of DOR+. Individual discriminatory FIRM score of each diagnostic criterion (idFIRM\text{c1 to n}) for FIRM\text{type 1 to n} was derived by converting its linear DOR+ value in to logarithmic value.

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Table 1: The clinical classification of total joint arthroplasty Failure by Immune Reaction to Metal ions (FIRM).

| FIRM\text{type1} | Presenting complication is predominantly dermatitis with minimal or no joint pain |
|------------------|----------------------------------------------------------------------------------|
| FIRM\text{c1a}  | Localized cutaneous reaction (either new or exacerbation of previous lesion; located around the replaced joint or elsewhere) |
| FIRM\text{c1b}  | Generalized cutaneous reaction (either new or exacerbation of previously existing lesions) |
| FIRM\text{type2} | Presenting complication is predominantly painful joint effusion (with negative work-up of infection or neoplasm) with minimal or no dermatitis |
| FIRM\text{c2a}  | Well fixed implants without evidence of osteolysis/cystic lesions |
| FIRM\text{c2b}  | Well fixed implants with evidence of osteolysis/cystic lesions |
| FIRM\text{c2c}  | Aseptic loosening of component(s) |
| FIRM\text{c2d}  | Instability/Dislocation of the joint |

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Figure 1: PRISMA Diagram showing the study flow from identification of original articles through meta-analysis.
If all the individual diagnostic criteria identified for FIRM type 1 to n are perfectly independent of each other, the total discriminatory FIRM score of FIRM type 1 to n that would determine the treatment threshold for revision surgery can be mathematically be expressed as,

$$td_{FIRM_{type\ 1\ to\ n}} = id_{FIRM_{c1}} + id_{FIRM_{c2}} \ldots \ldots + id_{FIRM_{cn}}$$

Where,

- $td_{FIRM_{type\ 1\ to\ n}}$ is the total discriminatory score of FIRM type 1 to n (in logarithmic scale),
- $id_{FIRM_{c1}}$ is the individual discriminatory FIRM score of a diagnostic criteria $c_1$ for FIRM type 1 to n (in logarithmic scale), and so on.

If we found that the individual diagnostic criteria identified for FIRM type 1 to n were partially independent of each other, we decided to fix the treatment threshold for revision surgery at 70% of the calculated $td_{FIRM_{type\ 1\ to\ n}}$ for FIRM type 1 to n.

### Results

When Pub Med and Embase were searched as per method 1 using the term ‘Metal Allergy Joint Replacement’ without applying any other filters, 195 and 76 articles were identified respectively. When a similar search was conducted as per method 2 using the term ‘Metal Hypersensitivity Joint Replacement’, 178 and 194 articles were identified in Pub Med and Embase respectively. The term that yielded maximum number of results among the two search terms was selected for record identification and abstract review. After removing the duplicates there were 334 articles available for screening of abstracts (Figure 1). When we included our unpublished record (MPN), we had a total of 335 records available for screening. We excluded 271 records among the final 335 that were screened, as they were not relevant to answer this study objective. We full text reviewed the remaining 68 records among the 335 records that were screened. Out of 68 records that were full text reviewed, 29 records were excluded for various reasons. Hence we identified a total of 39 original articles with data relevant to this study. During the process of full text review of these 39 records, we could not identify any additional original articles that were not previously identified by either Pub Med or

### Table 3: The diagnostic scoring system for FIRM type 1 is represented below.

| Diagnostic criteria(C1-8) of FIRM type 1 | idFIRM scores |
|----------------------------------------|--------------|
| C1. Histopathology of skin/regional lymph node(s) | 0.95 |
| Positive | 0.95 |
| Negative | 0 |
| C2. Serum (Co>19µg/L or Cr>17µg/L) | 0.95 |
| Positive | 0.95 |
| Negative | 0 |
| C3. Skin patch test (Ni or Co or Cr) | 0.80 |
| Positive | 0.80 |
| Negative | 0 |
| C4. Sex | | |
| Female | 0.05 |
| Male | 0 |
| C5. Joint | | |
| Hip | 0.43 |
| Knee | 0 |
| C6. Articulation | | |
| M type | 0.43 |
| N type | 0 |
| C7. Imaging (XR/CT/MRI) | | |
| Positive | 0.43 |
| Negative | 0 |
| C8. LTT/LAT/LFT | | |
| Positive | 0.26 |
| Negative | 0 |

The recommended cutoff value for the treatment threshold is 3.38.
Embase search but had the clinical data on FIRM in patients with TJA. Hence, we had a total of 39 articles available for meta-analysis.

**Part I: FIRM clinical classification**

Based on predominant symptom of clinical presentation, we classified FIRM into two types i.e., FIRM_{type1} and FIRM_{type2} (Table 1).

Further subclassification of cutaneous reaction was attempted based on the location of skin lesions i.e., localized or generalized. Similarly, subclassification of painful joint effusions was attempted based on further clinical qualifiers i.e., evidence of osteolysis or component loosening or dislocation.

**Part II: FIRM diagnostic scoring system**

Nine among the 39 articles selected for final analysis had data on FIRM_{type1} (Table 2). A total of 12 joints that developed FIRM type1 were identified among these 9 articles. We identified 8 individual

| First Author (alphabetical order) | Sex | Joint | Articulation | Patch test (Co, Cr, Ni) | LTT/LAT | HP | Imaging (XR/CT/MRI) |
|-----------------------------------|-----|-------|--------------|------------------------|---------|----|---------------------|
| Anand A [14]                     | ♂ 1 | THA   | M            | +                      | +       | 1  | 0                   |
| Berghs Schmidt P [15]            | ♂ 1 | THA   | M            | +                      | +       | 1  | 0                   |
| Bernasek TL [16]                 | ND  | 0     | 1            | 1                      | +       | 1  | 0                   |
| Blumenfeld TJ [17]               | ♂ 1 | 0     | 0            | +                      | +       | 1  | 0                   |
| Campbell P [18]                  | 4   | 1     | 0            | +                      | +       | 1  | 0                   |
| Carr AM [19]                     | 2   | 1     | 0            | +                      | +       | 1  | 0                   |
| Engh CA Jr [20]                  | ND  | 1     | 0            | +                      | +       | 1  | 0                   |
| Jensen P [21]                    | ♂ 1 | 0     | 1            | +                      | +       | 1  | 0                   |
| Kemp MA [22]                     | 3   | 0     | 2            | +                      | +       | 1  | 0                   |
| Kiran M [23]                     | ♂ 1 | ND    | M            | +                      | +       | 1  | 0                   |
| Korovessis P [2]                 | ND  | 0     | 0            | +                      | +       | 1  | 0                   |
| Kosukegawa [24]                  | 1   | 0     | 0            | +                      | +       | 1  | 0                   |
| Kumar [25]                       | 1   | 0     | 0            | +                      | +       | 1  | 0                   |
| Lohmann CH [26]                  | 24  | 3     | 0            | +                      | +       | 1  | 0                   |
| Mahendra G [27]                  | ND  | 0     | 0            | +                      | +       | 1  | 0                   |
| Melosev I [28]                   | ND  | 25    | 0            | +                      | +       | 1  | 0                   |
| Mikael MM [29]                   | 0   | 2     | 0            | +                      | +       | 1  | 0                   |
| Mc Master WC [30]                | 1   | 0     | 0            | +                      | +       | 1  | 0                   |
| Nett MP [31]                     | 0   | 2     | 0            | +                      | +       | 1  | 0                   |
| Pandit H [32]                    | 13  | 0     | 13           | +                      | +       | 1  | 0                   |
| Perumal V [33]                   | 1   | 0     | 0            | +                      | +       | 1  | 0                   |
| Rajpara A [34]                   | 5   | 13    | 0            | +                      | +       | 1  | 0                   |
| Roessler PP [35]                 | 0   | 1     | 0            | +                      | +       | 1  | 0                   |
| Singh C [36]                     | 1   | 0     | 1            | +                      | +       | 1  | 0                   |
| Thakur RR [37]                   | 4   | 1     | 0            | +                      | +       | 1  | 0                   |
| Theruvil B [38]                  | 3   | 0     | 3            | +                      | +       | 1  | 0                   |
| Thomas P [39]                    | 8   | 16    | 0            | +                      | +       | 1  | 0                   |
| Toms AP [40]                     | 0   | 1     | 0            | +                      | +       | 1  | 0                   |
| Vivegananathan B [31]            | 1   | 0     | 0            | +                      | +       | 1  | 0                   |
| Willert HG [41]                  | 9   | 18    | 0            | +                      | +       | 1  | 0                   |
| DOR+                             | 2.3 | 0.36  | 0.36         | 0.36                   | 0.36    | 0  | 0.17                |
| idFIRM_{type1} score             | 1.32| 1.32  | 1.32         | 1.32                   | 1.32    | 1.32| 0.91                |

**Table 4:** The results of meta-analysis of FIRM_{type2} diagnostic criteria.

LTT/LAT/LFT: Lymphocyte Transformation Test(s)/Lymphocyte Activation Test(s)/Lymphocyte Function Test(s); HP: Histopathology of joint tissue; XR/CT/MRI: X-rays/Computerized Tomography/Magnetic Resonance Imaging; THA: Total Hip Arthroplasty TKA: Total Knee Arthroplasty; M: Hip or knee implant with Metal on Metal bearings or implants with Mores taper between two modular CoCr components; N: Hip or knee implant with CoCr component but has no Metal on Metal bearings or Mores taper between two modular CoCr components; NT: Not Tested; ND: No Data; Of the three reported cases, one case not included because of dual diagnosis of infection and immune reaction to metal ions; @: Unpublished data of author (MPN); #: Histopathology finding were not reported in 2 among the total of 19 patients.
diagnostic criteria that were at least partially independent of each other. When we tested the statistical weight of these 8 individual diagnostic criteria, we identified that all are useful for diagnosis of FIRM type2; but the contribution of each to the identified FIRM type2 idFIRMc1 to 7 scores to tdFIRM type2 idFIRM score was variable. The idFIRMc1 to 7 scores of FIRM type2 were found to be useful for diagnostic purpose. The contribution of each to the identified diagnostic criteria were partially independent, the treatment threshold for revision surgery was estimated to be 3.39 i.e., 70% of 4.85.

**Discussion**

**Significance of the study**

This is the first proposed comprehensive classification system of FIRM in patients with either THA or TKA that identified and statistically weighed the strength of individual diagnostic criteria, developed of a clinically applicable diagnostic scoring system and estimated a threshold level for the revision surgery.

**Concerns**

There are three specific concerns that we should address in detail regarding the methodology used in this meta-analysis. First, is the use of DOR+ (also called Likelihood ratio) as a single indicator of a diagnostic criterion’s performance compared to the traditional paired indices of test validity such as predictive values. The attractive features of likelihood ratios that are not shared by the predictive values are applicability to specific patient rather than relate test results to populations, applicability across the disease frequencies and ability to combine the tests in order to refine the clinical judgment [40].

Second issue concerning the methodology is to provide an explanation for use of logarithmic scale than a more familiar linear scale. The best way to estimate the strength of a differentially used diagnostic criteria across the studies is to convert linear scores to logarithmic scores so that, the plotted score-treatment threshold curve will be changed from hyperbolic to sigmoid curve where usually, between 25% to 75% of the maximum response, the relation between the diagnostic scoring system and the treatment threshold will be linear, so that a better understanding and interpretation is possible. In other words, if all the included studies for meta-analysis did used exactly the same set of diagnostic criteria for decision making in all their cases, there would have been no need to convert the linear scores into logarithmic scale.

Third and the most important issue concerning this scoring system is the selection of 70% of tdFIRM score as the treatment threshold. Nomogram created for the Basey theorem indicates that an odds ratio of 10 to 1, 100 to 1 and 1000 to 1 are equivalent to a post-test probability of 50%, 75% and 99% respectively [41]. As the minimum required tdFIRM type1 and tdFIRM type2 score for thresholding is considered 3.38 and 3.39 respectively, the odds of being correct is more than 2000 to 1 or a post-test probability of approximately 99%.

**Clinical application**

It is important to understand that though 7 out of 8 tested diagnostic criteria for FIRM type1 and FIRM type2 are the same, their contribution to their respective tdFIRM score is different based on the strength of the odds ratio of each individual diagnostic criteria. The gender of the patient, the joint under evaluation and availability of previous surgical records provides pre-test tdFIRM score for a patient under evaluation for FIRM. Depending on the pre-test tdFIRM score, this scoring system provides clinicians with flexibility to decide which among the 5 tests (Histopathology/Imaging/Path test/Serum...
Table 6: Etiology and pathogenesis of FIRM in patients with total joint arthroplasty.

| Principle stimulus | Type1 FIRM | Type2 FIRM |
|--------------------|------------|------------|
| Particle size      | Sub micrometer | Sub nanometer | Sub micrometer |
| Joint tissue metal content | Not known | ↑ | ↑ |
| Joint metal ion level | Not known | ↑↑ | ↑ |
| Serum metal ion level | ↑↑ | ↑ | ↑ |
| Interaction with APC | Extra cellular | Intracellular | Extra cellular |
| Pathway | $T_{eff}$ inflammation | $T_{eff}$ inflammation | $T_{eff}$ inflammation |
| Helper T cells | $T_{eff}$ cell | $T_{eff}$ cell | $T_{eff}$ cell |
| Cytokine mediator | IL-17 | INF-$\gamma$ | IL-17 |
| Effector cells | Activated CTLs | Activated Macrophage | Activated CTLs |
| Histological appearance | PVLI, DLI | Coagulative necrosis | PVLI, DLI |
| Clinical effect | Synovitis (=pain/effusion) | Dermatitis | Osteolysis (= pseudo tumors, aseptic loosening) | Synovitis (=pain/effusion) |

Conclusion

This meta-analysis provides a clinically useful tool for decision making when evaluating a patient suspected to have developed clinical complication by immune reaction to metal debris from arthroplasty. Future studies on FIRM should utilize this scoring system in decision making and critically evaluate its validity.

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