Introduction

Fibromuscular dysplasia (FMD) is a rare noninflammatory and nonatherosclerotic vascular disease that mainly affects the renal arteries, but less commonly the carotids and vertebral arteries (1-3). FMD was first described in 1938 by Leadbetter and Burkland (4), and it was later classified pathologically, by Harrison and MacKormack in 1971 (5). The most common type of FMD is the “medial fibroplasia” (80-90%) with its characteristic picture of the “string of beads” pattern. Less common types are the “intimal” and “adventitial” FMD which are much less common accounting for 10% and <5% of cases, respectively. The prevalence of FMD in the general population is not well known. Estimates are derived from screening kidney donors, with a prevalence of about 2.6%. Among patients with renovascular hypertension (RVH), its incidence is about 10%, whereas 80-90% of RVH is due to atherosclerotic renal artery stenosis (ARAS). The treatment of choice of hypertension due to FMD is percutaneous renal angioplasty (PTRA). In contrast, hypertension due to ARAS is not frequently responsive to PTRA. In order to achieve successful control of hypertension in patients with FMD, a combination of PTRA with drugs that block the renin-angiotensin-aldosterone system (RAAS) is often necessary. The purpose of this review was to search the literature for newer diagnostic methods and treatment of FMD. Therefore, a Medline search of the English literature of published papers between 2008 and December 2013 was performed. Of 58 papers reviewed, 19 pertinent papers were selected including, studies, reviews, registries and case reports. The information from these studies together with collateral literature will be discussed in this concise review.

Keywords: Renovascular hypertension (RVH); fibromuscular dysplasia (FMD); medical treatment; angioplasty
this review we performed a Medline search of the English language literature using the terms RAS and FMD of the renal arteries from January 2008 to December 31 2013. From the 58 papers reviewed, 19 pertinent papers were selected. The information from these papers together with collateral literature will be discussed in this concise review.

Pathophysiology of FMD

FMD of the renal arteries is a noninflammatory vascular disease, which commonly affects the renal, carotid, and vertebral arteries. However, FMD can also affect arteries in other vascular territories. As described above, renal artery FMD (RAFMD) typically presents anatomically in three different types, including “medial”, “intimal”, and “adventitial” fibroplasia (3).

Medial FMD

This type is the most common accounting for 80-90% of the cases. It presents with its classic appearance the “string of beads” due to the alternating areas of constrictions and post-stenotic dilations of the renal artery (Figure 1). The lesions affect mainly the medium or distal third of the main renal artery, but they may extend into the proximal portion of the renal artery branches. This lesion can be bilateral in 60% of the cases.

Intimal FMD

This is the second most common (10%) presentation of FMD. It is due to collagen deposition within the intimal complicated often by a fragmented or duplicated internal elastic lamina. Angiographically, it is distinct from the medial fibroplasia because the intimal fibroplasias cause a focal fibrotic constriction that results in a concentric stenosis or long tubular lesion.

Adventitial FMD

This is the least common (<5%) presentation of FMD. It is usually due to the hypertrophy of connective tissue at the junction of the medial and adventitial layers of the renal artery (3). These lesions cause unifocal FMD, which can be found at the ostium, the trunk, or the bifurcation of the renal arteries.

RAFMD affects mostly young women in the prime of their life. The most common clinical symptoms are severe hypertension, headaches, pulsatile tinnitus, and dizziness. The pathophysiologic cause of hypertension in unilateral RAFMD is activation of the renin-angiotensin-aldosterone system (RAAS) secondary to postenotic drop of renal artery pressure and renal ischemia. The increased systemic BP leads to pressure diuresis from the contralateral unobstructed kidney causing plasma volume contraction and further stimulation of RAAS. In these cases, relieve of the obstruction leads to prompt reduction of the activity of RAAS and decrease in BP. In bilateral RAFMD, there is activation of RAAS but the kidneys are not subjected to increased systemic BP and therefore, do not respond with pressure diuresis. In these patients, the hypertension is due to a combination of volume expansion and RAAS activation, and bilateral angioplasty with relief of obstruction in both kidneys will lead to prompt reduction of the activity of RAAS and BP. In contrast to atherosclerotic renal artery stenosis (ARAS), RAFMD almost never, leads to complete obstruction of the renal artery.

Extrarenal manifestations of FMD

Although FMD mainly affects the renal arteries, it can less frequently affect arteries in other vascular territories, as described in the following case studies.

Carotid arteries

Medial FMD (string of beads) of the right internal carotid
(ICA) was found in a 52-year-old man who was diagnosed with an acute right posterior parietal brain infarct. This infarct was due to complete occlusion of right ICA from a dissection. The patient was treated successfully with aspirin and clopidogrel without further sequelae (9).

In another case, medial FMD of both carotid arteries was diagnosed in a 52-year-old female, who presented with severe hypertension, severe headaches, right ocular pain, diplopia and amaurosis fugax (10). Both renal arteries were normal. She was treated conservatively with antihypertensive and antiplatelet (aspirin) drugs and had no symptoms one year later.

Brachial arteries
Medial FMD of the left brachial artery was diagnosed in an 83-year-old man with end stage renal disease causing access problems for hemodialysis. The brachial stenosis was treated successfully with balloon angioplasty (11).

Ulnar and radial arteries
Medial FMD was diagnosed in a 20-year-old male who presented with ischemia and cyanosis of the right hand of six weeks duration. Examination revealed significant stenosis of the right ulnar artery, but the radial artery was not significantly narrowed. He was successfully treated with excision of the diseased segment of the ulnar artery with end to end anastomosis (12).

Mesenteric arteries
A rare case of periaortic FMD of the right mesenteric artery was diagnosed in an elderly patient complaining of abdominal pain, constipation, and bowel distention. He was treated successfully with a right hemicolectomy. The rest of the colon and cecum were normal (13).

External iliac and femoropopliteal arteries
Two cases of medial FMD with obstruction of the iliac and femoropopliteal arteries were diagnosed. In one case a 63-year-old woman presented with disabling left leg claudication (ABI 0.37) and hypertension. She was diagnosed with medial FMD with bilateral common iliac artery aneurysms, and alternating aneurysms and stenoses of both popliteal and superficial femoral arteries. She was treated successfully with resection of the aneurysms and reconstruction of the superficial femoral arteries (14).

The second case concerns a 37-year-old man presented with sudden onset of back pain. He was diagnosed with acute pancreatitis and thrombus with dissection of the celiac trunk and right external iliac artery. He was treated successfully for his pancreatitis and with a transluminal coil embolization of the iliac trunk.

Diagnostic evaluation of RAFMD
Catheter-based angiography remains the gold standard for the accurate diagnosis of RAFMD, because it can visualize the main renal arteries as well as the smaller branch vessels. In addition, catheter-based angiography has the advantage that a pressure wire and intravascular ultrasound (IVUS) imaging can be used to visualize the fibrous rings and webs that cause the intravascular obstruction and increase the pressure gradient (Figure 2), as well as evaluate the success of PTRA in cracking the rings (15-17). Other diagnostic modalities like CT angiography (CTA) and contrast-enhanced magnetic resonance angiography (MRA) display good specificity in the detection of RAFMD, and are thus recommended as imaging modalities for its diagnosis (18). In a study by Willoteaux et al. (18), in patients with RAFMD the sensitivity and specificity of contrast-enhanced MRA were 97% and 93%, respectively for the accurate diagnosis of main renal artery disease and it was highly correlated with digital subtraction angiography. In addition to these...
diagnostic methods, the measurement of translesional systolic blood pressure (SBP) gradient during catheter angiography should be measured as well. A systolic SBP gradient ≥20 mmHg is considered diagnostic for significant RAS (19). Another noninvasive method to evaluate the severity of RAS is the use of duplex scan (20). With this method the peak systolic velocity (PSV) in the stenosed renal artery can be measured. From this measurement, the ratio of PSV of the renal artery to PSV of the aorta (RAR) can be determined. The diagnostic values for significant RAS from this test are listed in Table 1.

**Table 1 Classification of renal artery stenosis by duplex scanning of the renal artery**

<table>
<thead>
<tr>
<th>Renal artery diameter reduction</th>
<th>PSV, cm/sec</th>
<th>RAR</th>
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<tbody>
<tr>
<td>Normal</td>
<td>&lt;180</td>
<td>&lt;3.5</td>
</tr>
<tr>
<td>&lt;60%</td>
<td>≥180</td>
<td>&lt;3.5</td>
</tr>
<tr>
<td>≥60%</td>
<td>&gt;180</td>
<td>≥3.5</td>
</tr>
<tr>
<td>Occlusion</td>
<td>No signal</td>
<td>No signal</td>
</tr>
</tbody>
</table>

PSV, peak systolic velocity; RAR, renal aortic ratio. Adapted with permission from Caps et al. (20).

**Table 2 clinical factors favoring medical therapy or revascularization of RAS**

<table>
<thead>
<tr>
<th>Medical therapy and surveillance</th>
<th>Revascularization ± medical therapy</th>
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<tbody>
<tr>
<td>Controlled BP with stable renal function</td>
<td>Progressive decline in GFR during treatment of hypertension</td>
</tr>
<tr>
<td>Advanced age or limited life expectancy</td>
<td>Failure to achieve adequate BP control with optimal medical therapy</td>
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<tr>
<td>Extensive comorbidities that make revascularization risky</td>
<td>Rapid or recurrent decline in GFR in association with BP reduction</td>
</tr>
<tr>
<td>High risk for atheroembolic complications</td>
<td>Decline in GFR with treatment with ACE inhibitors or ARBs</td>
</tr>
<tr>
<td>Concomitant renal parenchymal disease (interstitial nephritis, diabetic nephropathy etc.)</td>
<td>Recurrent CHF in patients with adequate left ventricular function</td>
</tr>
</tbody>
</table>

Abbreviations: RAS, renal artery stenosis; BP, blood pressure; ARB, angiotensin receptor blocker; RAS, Renal artery stenosis.

**Treatment of patients with RAFMD**

The treatment of choice in patients with RAFMD with hypertension is PTRA, with or without stent placement. This leads to successful control of BP either alone or less frequently in combination with pharmacological antihypertensive treatment. In contrast, in patients with ARAS, PTRA is less frequently successful in lowering the BP and will often require the addition of medical therapy. Another difference between RAFMD and ARAS is that RAFMD almost never causes renal functional impairment while ARAS is very frequently associated with hypertension and renal functional impairment. The American Heart Association has issued guidelines for the treatment of ARAS, which may help guide therapeutic approaches for patients with RAFMD (21). These guidelines regarding initial medical or interventional therapy are summarized in Table 2.

The following paragraphs summarize findings from studies evaluating the effectiveness and safety of PTRA in patients with RAFMD and hypertension either alone or in combination with antihypertensive drugs.

Øvrehus et al. (22), retrospectively analyzed the results of 12 patients with FMD and hypertension who were treated with PTRA and followed for an average of 13 years. In five of 12 (41.7%) the BP was normalized without medical therapy. Their BP dropped from a baseline of 170/100 to 130/78 mmHg after the intervention and the number of antihypertensive drugs was decreased from 2.7 to 0. In the other seven (58.3%) the BP dropped from a baseline 177/99 to 153/90 mmHg and the number of drugs was decreased from 3.6 to 1.0. Their BP and treatment remained the same for the whole follow-up period of 1,000 days (33.3 months). There was no change in the serum creatinine values in both groups. This study does not provide any information regarding the sex and age of patients.

Davies et al. (23), also analyzed retrospectively the results of 29 female patients with RAFMD, mean age 45 years (range, 18-80 years), who were treated with PTRA ± stent between January 1999 and December 2007. The baseline BP in the 21 (72.4%) of responders to PTRA was 169±20/92±13 mmHg, and in the eight (27.6%) nonresponders to PTRA the baseline BP was 180±36/91±14 mmHg. History of hypertension ≤8 years was present in 86% of the patients. PTRA without stent was initially used in all patients and stent was used only in cases of non successful PTRA or complications from the procedure. The patency rates with primary and assisted PTRA were 66% and 87%, respectively at five years. The hypertension...
was either improved or cured in 72% and 71% of patients, respectively at three months and five years after the procedure. The predictors of long-term success were the duration of hypertension ≤8 years, eGFR ≥60 mL/min/1.73 m², ipsilateral kidney size ≥9 cm, fasting blood glucose <110 mg/dL, and triglycerides <150 mg/dL.

Barrier et al. (24), also analyzed retrospectively, the findings of 30 patients with the less common types of FMD of unifocal and nonmedial dysplastic renal arteries, who were treated between 1994 and 2006. Of the 30 patients, 24 (80%) were females and six (20%) were males mean age 29.6 years (range, 13-51 years). Hypertension was the main presentation for treatment in 28/30 patients. Of the other two, one was treated for worsening of preexisting renal failure and the other for renal protection of a solitary kidney. The renal function of the 28 patients was within normal limits. All patients were treated initially with PTRA and stent was used only, when PTRA was not successful (seven patients). The immediate procedural success rate was 65% and increased to 82% with the additional procedure. Procedural complications occurred in 30%, but they were not serious and all were managed successfully. The initial success rate with improved or cured hypertension was 92%, but hypertension recurred later in 42% of patients. After the second procedure the BP benefit increased to 78% and was maintained for 84 months.

Thatipelli et al. (25), also made a retrospective analysis of the data from 16 older patients treated with PTRA for RAFMD and hypertension between 1999 and 2005. The mean age of the patients was 66±11 years and 14 (88%) of the patients were women. The median follow-up period was 21 months (range, 1.03-85.77 months). The duration of hypertension ranged from two to 24 years and 13 patients had developed severe, resistant to treatment hypertension. Two patients had chronic renal failure with serum creatinine levels 1.5 and 2.4 mg/dL, respectively. A total 21 procedures were performed, 18 on the right renal artery, one on the left renal artery and two were bilateral. Of the 21 procedures, 18 (95%) failed 12.8 months later. The main reason for the failure was prolonged duration of hypertension in 14 (93%) of patients. The mean number of antihypertensive medications before and after PTRA was 2.75±1.25 and 3.0±1.24, respectively (ns). The percentage of failures for BP control at 1, 6, and 12 months was 42%, 58%, and 79%, respectively. This study highlights the poor-, short- and long-term hypertension success rates of PTRA in older patients with RAFMD and hypertension.

Mousa et al. (15), also analyzed the data from 35 patients with RAFMD, of whom 32 (91%) were women. The mean age of the patients was 61.9 years and the main reason for treatment was sustained hypertension. In these patients 43 procedures were performed on 43 renal lesions and the initial success rate was 100%. Of the 35 patients, 27 had isolated RAFMD, 4 had RAFMD with atherosclerosis in the same artery, three with atherosclerosis in the opposite artery and 17 (48.6%) had bilateral RAFMD. The short-term reduction of SBP with PTRA was from (mean ± SEM) 162.2±3.8 to 132.6±2.4 mmHg (P<0.001), and for the diastolic BP (DBP) from 83.4±2.3 to 65.5±3.1 mmHg (P<0.001). The long-term (4.8±0.5 years) reduction in SBP was 142.6±3.6 mmHg from baseline (P<0.001), and in DBP 76.5±2.4 mmHg from baseline (P=0.02). Also, the number of antihypertensive medications was decreased from a baseline of 3.1±0.24 to 2.7±0.27 (P=0.03). Freedom from recurrent hypertension at one, two, and three years was 93%, 75%, and 41%, respectively. The eGFR increased with treatment in 26 patients, from a baseline of 71.1 to 75.6 mL/min (P=0.354). In ten patients with baseline eGFR of <60 mL/min, it increased to >60 mL/min in only two patients. The long-term patency rate was 95%, 71%, and 50% at one, five, and nine years, respectively, whereas the assisted primary patency rate remained at 100% for the nine years of follow-up. This study also, demonstrates the poor long-term response of BP to treatment in older patients.

Trinquart et al. (26) examined BP control and complication rate from PTRA in patients with RAFMD in a review and meta-analysis of 47 studies comprising 1,616 patients. Their baseline clinical characteristics are listed in Table 3. There was a great heterogeneity among the

| Table 3 Baseline characteristics of patients with hypertension due to RAFMD treated with PTRA |
|-----------------------------------------------|----------|----------|---------|
| Age (years)                                  | Studies (No) | Patients (No) | Mean |
| Men                                          | 34        | 974       | 42     |
| Bilateral RAFMD                              | 31        | 913       | 19%    |
| BP (mmHg)                                    | 30        | 873       | 26%    |
| Antihypertensives (No)                       | 20        | 605       | 174/108|
| Duration of HTN (years)                      | 12        | 486       | 2.02   |
| Medial RAFMD                                 | 13        | 442       | 7.3    |
| Table constructed from data by Trinquart et al. (26). Abbreviations: RAFMD, renal artery fibromuscular dysplasia; PTRA, percutaneous renal angioplasty. | 7         | 213       | 67%    |

Abbreviations: RAFMD, renal artery fibromuscular dysplasia; PTRA, percutaneous renal angioplasty.
RAFMD is a fairly uncommon disease and a rare cause of hypertension. Its prevalence in the general population is not currently known, but it is estimated from populations undergoing screening for potential kidney donation. In a recent study that evaluated 2,640 living kidney donors free of hypertension at the time of screening, 68 (2.6%) had RAFMD (7), whereas in a previous study of 1,862 subjects evaluated as potential kidney donors, 3.8% (71/1,862) had RAFMD (27). In this study as well the mean age of the subjects was 50.8 years and 75% were females. However, among patients with renovascular hypertension (RVH), the incidence of RAFMD is about 10%, since the greatest majority (80-90%) of patients have ARAS (28). RAFMD is a disease that affects mostly females in the prime of their lives, although it can occur at any age. Of 68 subjects identified with RAFMD among the 2,640 subjects screened for kidney donors, 59 (86.8%) were women mean age 52±10 years (7). Also, in the US Registry of 447 patients with RAFMD, 91% were women mean age 51.9±13.4 years (8). The disease has also been described in children and adolescents 2-18 years old (29). RAFMD is a noninflammatory fibrodysplastic condition affecting primarily the renal arteries, but also the carotid and vertebral arteries as well as other arteries of the body (9-14). Its etiology is not known at present, and its hereditary nature is disputable, although it has been found in close relatives of patients with RAFMD. In a retrospective analysis of 104 patients with RAFMD, 11% had a history of familial disease (30). Cigarette smoking has also been implicated as a possible cause of RAFMD (31). In a study of 337 French patients with RAFMD, the proportion of smokers was 30% compared to 18% for non smokers (P<0.001). The most common clinical presentations of patients with RAFMD are severe, uncontrolled hypertension, pulsatile tinnitus and dizziness. Renal failure is a rare symptom of RAFMD and only occurs with complications of the disease like renal artery dissections or thrombosis due to aneurysms in bilateral RAFMD (32). In contrast in patients with ARAS, renal failure is a common presentation (28). Another difference between RAFMD and ARAS is that PTRA ± stent is the treatment of choice for RAFMD, whereas this modality is not always successful in patients with ARAS and most studies have shown no difference between interventional and medical therapy in lowering the BP (19,28). In addition, PTRA in patients with RAFMD is not associated with atheroembolic complications of the kidneys in contrast to ARAS where these complications are frequent due to atherosclerotic disease and the overall prognosis is more favorable in patients with RAFMD. With respect to BP response to treatment, the age and duration of hypertension are very important. In a recent review and meta-analysis, the age and the duration of hypertension were inversely associated with the BP response to treatment (26). In contrast, the response or cure of hypertension after PTRA in children is very high because the age and duration of hypertension are much smaller (29). It is quite possible that long-standing hypertension in patients with RAFMD could damage the opposite unprotected kidney leading to sustained hypertension, which could become unresponsive to PTRA. Therefore, the early diagnosis and treatment of patients with RAFMD is very critical. In addition to

<table>
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<th>Table 4 Percentage of blood pressure response to PTRA by different definitions in patients with FMD of the renal arteries</th>
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<td></td>
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<tr>
<td></td>
</tr>
<tr>
<td>Any BP Resp [%]</td>
</tr>
<tr>
<td>BP &lt;140/90 [%]</td>
</tr>
<tr>
<td>DBP &lt;90 [%]</td>
</tr>
</tbody>
</table>

All these BP responses to PTRA are without treatment. Resp, response. Constructed from data by Trinquart et al. (26).

Abbreviations: FMD, fibromuscular dysplasia; PTRA, percutaneous renal angioplasty.
age and long-standing hypertension, the presence of risk factors such as diabetes mellitus, dyslipidemia, and renal insufficiency are contributing factors to the poor response of BP to treatment and these risk factors should be treated together with the treatment of hypertension. The treatment of hypertension due to RAFMD can be either medical, interventional, or the combination of both. The drugs of choice are those that block the RAAS, such as angiotensin converting enzyme inhibitors (ACEI), angiotensin receptor blockers (ARB), and direct renin inhibitors (DRI) since this type of hypertension is renin dependent. In addition, RAAS blockers can be used in combination with diuretics, calcium channel blockers (CCB) and beta-blockers (b-blockers). Occasionally, in severe bilateral RAFMD the use of RAAS blockers could lead to significant increase in serum creatinine and BUN due to drop in glomerular filtration pressure from the decrease in the efferent arterial pressure produced by these drugs (33). This complication is completely reversible with the discontinuation of the medications. Another potential modality of treatment in patients with RAFMD is renal denervation. Recently, a case of renal denervation was reported in a 62-year-old female with severe, resistant to medical therapy hypertension. Renal denervation resulted in additional BP reduction to the one achieved by the baseline medical therapy (34). However, large experience with the use of denervation for the treatment of patients with hypertension due to RAS is not available because these patients were excluded from participation in the large denervation studies. Unfortunately, the largest denervation study SIMPLICITY 3 (35), was recently stopped prematurely, because the BP reduction was not better than placebo (sham denervation). Therefore, the results of this study will, most likely, put in jeopardy the future use of renal denervation for the treatment of resistant to treatment hypertension.

Acknowledgements

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