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1 Takayasu Arteritis in Pregnancy: A Rare Case Report

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4

5 **Abstract**

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7 **Index terms—**
8 Takayasu Arteritis in Pregnancy: A Rare Case Report lata Assudani

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10 **2 I. INTRODUCTION**

11 Takayasu arteritis (TA), also known as "young female arteritis", is a rare and chronic inflammatory disease of
12 large vessels. The disease mainly affects women of reproductive age and Asian origin (1) . Moreover, TA leads to
13 several complications including occlusion as well as aneurysm formation in systemic and pulmonary arteries. Its
14 incidence is reported to be 13 cases per million population (2) . Pregnancy as such has no effect on the evolution
15 of the disease, however, its peak incidence is in second and third trimesters.

16 Thus, such patients warrant special attention during peripartum period owing to likelihood of development of
17 complications such as hypertension, multiple organ dysfunction and stenosis hindering regional flow leading to
18 restricted intrauterine fetal growth and low birth weight in babies (3)(4)(5).

19 Delay in diagnosis is quite common, so patients often conceive without prior knowledge of having TA or
20 having initiated specific treatment against it (6) Ideal management for pregnant patients with this disease still
21 poses a stringent challenge, an interdisciplinary collaboration of obstetricians, cardiologists, rheumatologists
22 and neurologists is often necessitated for an optimal maternal and fetal prognosis. Here is the case described
23 to enlighten the obstetricians on fetomaternal outcome and management of this infrequent but not uncommon
24 clinical entity encountered nowdays. Baby cried immediately after birth and was admitted in NICU in view of
25 Extremely preterm and very low birth weight of 1.1kg.

26 **3 II. CASE HISTORY**

27 Patient was shifted to CCU for post-operative monitoring which was uneventful. Interventional radiologist
28 reference done in view of definitive management of Takayasu arteritis and was advised PET CT and stenting
29 after funds are available.

30 Patient was discharged with baby on day 36 PNC.

31 **4 III. DISCUSSION**

32 Takayasu arteritis was first described in 1908 by 2 Japanese ophthalmologists, Mikito Takayasu and onishi, who
33 observed retinopathy in the absence of peripheral pulses. The cause is unknown, but it seems to be related to
34 autoimmunity, sex hormone (more common in young females) and genetics (demonstrated by the predisposition
35 of the human leukocyte antigen-HLABW52).

36 Disease progression typically occurs in various stages from acute inflammatory arteritis to lymphocytic
37 infiltration, intimal thickening, elastic tissue destruction, fibrosis and patchy minimal narrowing of arteries.

38 **5 London Journal of Medical and Health Research**

39 Depends on angiographic classification there are five types based on the involvement arteries (1,7) .

40 Type The incidence of Takayasu arteritis during childbearing years is relatively high, the management of
41 pregnancies with this disease is of great importance in clinical obstetrics. Pregnancy with Takayasu arteritis
42 can be complicated by hypertension, as seen in our case, and worsening of cardiovascular hemodynamic status.

43 Hypertension is a serious complication that can lead to intrauterine growth retardation, fetal hemorrhage, and
 44 maternal heart failure (8) . The increased intravascular volume seen during pregnancy may impair circulation
 45 and exacerbate maternal hypertension, aortic regurgitation, and congestive heart failure (9).

46 The disease causes various clinical conditions depending on the sites of constriction such as arm claudication,
 47 decreased arterial pulses, visual loss, stroke, aortic regurgitation, Hypertension, congestive cardiac failure.
 48 Hypertension is seen in 90% cases Takayasu arteritis. The clinical patterns of TA differ at the acute and
 49 chronic periods. In the acute period, systemic symptoms prevail, while in the chronic period, insidious ischemic-
 50 destructive signs are more prevalent.

51 These signs appear together with stenosis at a rate of 85%, dilatation at a rate of 2%, and stenosis and
 52 dilatation at a rate of 13% (10,11,12) .

53 The symptoms range from fever, fatigue, and weight loss to life-threatening hemoptysis and heart failure.

54 Diagnosis is usually based on clinical manifestations, inflammatory markers (acute phase reactants), and
 55 arteriography demonstrating aortic stenosis and of its branches.

56 Common features of active TA are fatigue, myalgia, arthralgia, and low-grade fever in initial stages and
 57 intermittent claudication, visual defects, and fainting attacks in later stages. Many may be diagnosed after
 58 clinical examination, when one or more peripheral pulses are not palpable or blood pressures vary in two limbs.

59 However, computed tomography or magnetic resonance angiography can detect TA even before the development of severe vascular compromise as in our case (13) B Recently, 18 FDG-PET scan has been added as an
 60 adjunct imaging modality in the armamentarium of rheumatologists and cardiologists to diagnose LVV, with a
 61 pooled sensitivity and specificity of 70.1% and 77.2%, respectively (14) . But this is currently not available in
 62 our hospital.

63 However, the gold standard for diagnosis still remains as vessel biopsy (10) which could not be performed in
 64 our case.

65 The management of TA is a multidisciplinary approach with the involvement of obstetricians, anesthesiologists,
 66 cardiologists, rheumatologists, and neonatologists. Ultimately, the aims encompass the control of inflammation,
 67 prevention, and treatment of complications like hypertension and occlusive or stenotic lesions (15) . The aims
 68 are control of inflammation, prevention and treatment of complications like hypertension and revascularization
 69 by percutaneous angioplasty, use of endoprosthesis, or surgical correction for occlusive and stenotic lesions.

70 When managing women of reproductive age with TA, preconception counseling is essential. In addition, such
 71 counseling will focus mainly on dosage adjustment, cessation of cytotoxic drugs, folic acid supplementation in
 72 the London Journal of Medical and Health Research

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75 Takayasu Arteritis in Pregnancy: A Rare Case Report periconception period, and the optimal timing of
 76 pregnancy. Similarly, the pregnancy should be ideally planned in remission phase and patients are encouraged to
 77 pursue an early booking for regular antenatal supervision. In addition to routine antenatal visits, serial monitoring
 78 of BP, renal function, cardiac status, and pre-eclamptic screening is vital in such patients. Furthermore, fetal
 79 surveillance is also necessary and will include daily fetal kick count, gravidogram, serial fetal biometry, biophysical
 80 profile, and fetal Doppler (16) Controlling BP during pregnancy may be difficult due to the physiological changes
 81 in this period.

82 Thus, any patient with TA should plan to conceive when the BP and disease are stable. It is also vital to adjust
 83 the antihypertensive medication and avoid angiotensin-converting enzyme inhibitors or angiotensin inhibitors.
 84 On the other hand, uncontrolled hypertension during pregnancy has been associated with abortion, stillbirths,
 85 aortic dissection, cardiac and renal insufficiency, stroke, and maternal death (17)(18)(19) .

86 Antihypertensive drugs and antiplatelets can be started as per need, as was in the present case. TA may
 87 respond symptomatically to corticosteroid therapy (first line drugs) at a dose of 1-2 mg/kg/bodyweight for 4
 88 weeks followed by slow tapering. However, chronic use of corticosteroids could lead to suppression of adrenal
 89 gland activity with inadequate release of endogenous corticosteroids in moments of stress, such as surgeries (20)
 90 . Also immune-suppressors including methotrexate and azathioprine are used.

91 Finally, vaginal delivery has proven to be the preferred mode of labor management for patients with TA.
 92 Additionally, epidural analgesia has been advocated for labor and delivery as well and delivery abbreviated
 93 by use of forceps. In our case, decision for emergency LSCS taken in view of USG S/O uteroplacental and
 94 fetoplacental insufficiency with previous lcs with short ICP and was uneventful.

95 Patient was monitored postoperatively in CCU and was transferred to medicine for further management, where
 96 patient continued on steroids as she was breast feeding and methotrexate was contraindicated and discharged
 97 with advise to monitor BP and follow up with BP charting after 3 months for revascularisation surgery. ¹ ²

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