Case Report of Diagnostic Evidence Depicting Palmaris Longus Origin Reversal: Ultrasound and MRI Evaluation

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Abstract

Introduction: The palmaris longus (PL) is very prone to anatomical variance. It may exhibit agenesis, reversal, and duplication. The reversal variant presents with the PL tendinous aspect proximally and the muscle belly distally. It is an important finding, and it is important to consider when making a differential diagnosis. This is the only reported case of a left distal forearm PL muscle variant with ultrasound and magnetic resonance imaging (MRI) in a Caucasian adult male.

Case Report: A 28-year-old Caucasian male presented to the clinic with an abnormal ultrasound finding of a left wrist mass. The patient was instructed to get an MRI, which demonstrated and further confirmed the reversed PL muscle.

Conclusion: Although rare, anatomic muscle variance may occur in many places of the body leading to symptoms that need attention. These variants should be considered in orthopedic differential diagnosis and ruled out appropriately with proper diagnostic techniques. By making the correct diagnosis, it leads to improved patient outcomes and satisfaction both conservatively and surgically.

Keywords: palmaris longus; anatomic variant; forearm musculature; forearm flexor.

Introduction

There are many cases of anatomical variance in the musculature of the upper extremity (UE). The fusiform palmaris longus (PL) muscle is prone to variance, and in order of prevalence; the completely absent is the most common, followed by the reverse, duplicated, bifid or hypertrophied form [1]. Opposite to normal anatomy, the reversed PL exhibits a proximal tendinous structure and a muscle belly distally [1]. A recent meta-analysis from 1975 to 2014 reported that there is only one medical record of magnetic resonance imaging (MRI) displaying the reversed PL [2]. The present report describes a case of a reversed PL diagnosed with ultrasound and MRI, with a literature review to demonstrate the vast spectrum of variation seen in the musculature of the UE.

Case Report

A right hand dominant, 28-year-old Caucasian male presented to the orthopedic associates of Southwest Ohio as a referral to a hand specialist due to an abnormal ultrasound finding and left wrist mass. The ultrasound of the suspected left wrist mass was executed 10 days prior, exhibiting increased echogenicity within the musculature of the left distal wrist at the site of the patient’s reported abnormality. This echogenicity was asymmetric when compared to the right wrist. The image of the left anterior distal forearm on ultrasound is displayed in Figure 1.

![ ultrasound image of wrist ]

Figure 1: Anterior left forearm muscle variant demonstrated on ultrasound imaging 1 month before the initial follow-up to presenting surgeon/ultrasound of the left anterior wrist area.
There was no evidence of a focal mass or fluid collection according to the ultrasound report. The patient presented complaining of left wrist swelling, with no pain. There was an unremarkable past medical, surgical, or family history significant to the chief complaint. As far as a differential, the questionable echogenicity within the musculature of the left distal forearm could have been related to a mild non-specific myositis. Moreover, there was no evidence of focal mass or fluid collection, so an MRI was considered to rule out a neoplasm. The patient completed the MRI in 10 days, and then followed up in a week to the clinic. The current status of the patient remained unchanged. The MRI of the patient demonstrated a palpable abnormality secondary to a congenital variation consisting of a PL muscle variant due to an accessory muscle belly volar to the flexor tendons. The signal intensity of this muscle appears within normal limits. This finding was confirmed on the retrospective review of the prior ultrasound. The abnormal muscle variant is displayed in the sagittal view T1 MRI in Figure 2, it is displayed in the coronal view T1 MRI in Figure 3, and it is displayed in the coronal view T2 MRI in Figure 4.

After consecutive diagnostic imaging techniques ruled out anything pathologic including neoplasm, the patient was counseled on the plan. Since the muscle variant was not painful and it did not limit functional outcomes, a non-operative conservative plan was instituted. The patient was informed to follow-up as needed, and if any concerning symptoms arose to schedule an appointment and new measures would be taken for continued care.

In the present case, the PL muscle belly originated at the distal forearm volar to the flexor digitorum profundus and superficialis tendons. The abnormality lied between the soft tissue markers placed at the site of the patient's complaint of the palpable abnormality.

**Discussion**

The standard anatomical origin of the PL is located on the medial epicondyle at the common flexor tendon, and the insertion is the palmar aponeurosis located volarly to the flexor retinaculum [1]. The PL tendon and the palmar aponeurosis are described as two separate entities based on their origin and development [3]. Neurovascularly, it is supplied by branches of the ulnar artery and innervated by branches of the median nerve [4].

This patient is a 28-year-old Caucasian male that presented with a left distal forearm PL muscle variant. Reimann et al. conducted the first major study that looked at 1600 UEs and the incidence of PL variations, and it showed that there were only three aberrancies of the attachment of the muscle in the 540 consecutive arms [5]. Moreover, Olewnik et al. created a classification system that described nine different types of anatomical presentations of the PL [6]. The patient’s abnormality presented with the muscle belly distally and tendinous portion proximally, which is described as the Type VI variant and is only identified in 2% of cases [6].
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The clinical significance of this case is placed on the nervous systems implications. Proximal muscle belly PL variants may predispose patients to median nerve compression [6]. Although it is much less common, compression of the ulnar nerve in Guyon’s canal has been reported as well [7]. Patients may also present with distal forearm edema and inflammation that may result in reduction of hand function due to wrist flexion movements [8].

It is important to understand anatomical variants of the PL and to incorporate into the differential diagnosis list because it plays a crucial role in reconstructive surgeries [9]. It is essential to clinically examine patients that present with an anterior distal forearm mass with suitable diagnostic techniques to avoid inappropriate surgical procedures.

Conclusion

This is a case report of a patient exhibiting symptoms from the reversed PL muscle in the forearm. Anatomical muscle variants should be considered in a differential diagnosis when there is an unknown muscular mass in a patient. Due to the minimal number of case reports on anatomical muscle variation, we suggest that with appropriate diagnostic techniques and proper management of such cases, it can lead to improved patient outcomes and satisfaction.

Clinical Message

With appropriate history and diagnostic techniques, considering anatomic muscle variants in your differential diagnosis can lead to improved patient outcomes and satisfaction.

References