Parry Romberg Syndrome: 7 Cases and Literature Review

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ABSTRACT

SUMMARY: Parry Romberg syndrome is a rare progressive hemiatrophy of the face that typically occurs in children and young adults and has a peculiar progression that ceases without apparent cause after a highly variable period. Only a subset of patients with Parry Romberg syndrome will develop secondary neurologic or ophthalmologic symptoms, and prognosis is highly variable. Inconsistency in the pattern of atrophy and the development of associated symptoms in patients with Parry Romberg syndrome has made it challenging to diagnose, prognosticate, and treat. The precise etiology of this disease remains unknown, but some authors have implicated sympathetic cervical ganglion dysfunction, abnormal embryogenesis, autoimmune and inflammatory mechanisms, or vasculopathy as potential causes. We present 7 cases of Parry Romberg syndrome and their associated clinical and imaging findings with specific attention to the radiographic characteristics of this disease.

ABBREVIATION: PRS — Parry Romberg syndrome

Initially described by Drs Caleb Hillier Parry and Moritz Heinrich Romberg in 1825 and 1846, respectively, Parry Romberg syndrome (PRS), also known as progressive facial hemiatrophy, is a rare, insidious, and self-limiting hemiatrophy of the face. It characteristically involves the skin and subcutaneous connective tissues and may later progress to affect the underlying musculature, cartilage, and osseous structures with or without development of neurologic symptoms. PRS typically presents initially in children and young adults and slowly progresses over a highly variable course ranging from 2 to 20 years, eventually reaching a “burned-out phase” and stabilizing for no apparent reason. This peculiar disease course, along with highly variable signs and symptoms, impedes consistent understanding of the underlying pathophysiology of PRS. Many theories about PRS have emerged throughout the years, attributing this syndrome to widely varying etiologies such as infection, trauma, sympathetic nervous system dysfunction, vascular abnormalities, inflammatory conditions, and autoimmune disorders, but at this time, a specific etiology remains uncertain.

Changes of cutaneous PRS may be subtle on imaging studies, but 20% of these patients will have intracranial manifestations that may not correspond to the severity of soft-tissue involvement or neurologic symptoms. Radiologic examinations may identify clinically occult intracranial involvement, facilitate the exclusion of other differential diagnostic considerations, and aid in monitoring disease progression. We present 7 cases of PRS to illustrate its wide variety of extracranial and intracranial imaging findings, along with a discussion of the clinical course, differential diagnoses, and treatment options. The role of the radiologist in diagnosis and management is emphasized.

Case Series

Case 1. A 44-year-old woman with known right-sided PRS underwent CT for the evaluation of disease extent and cosmetic surgical planning. CT demonstrated thinning of the skin and subcutaneous fat of the right side of the face extending inferiorly to the suprahyoid neck. There was atrophy of the right platysma and muscles of mastication, with substantial asymmetry in the size of the parotid gland (Fig 1A). Osseous asymmetry was most pronounced in the maxilla and zygoma, though the right hemimandible was smaller than the left one (Fig 1B). Considerable right enophthalmos was present (Fig 1C). No brain imaging was performed.

Case 2. A 14-year-old boy presented with progressive atrophy of the left side of the face. CT and MR imaging examinations revealed hemiatrophy most prominently involving left-sided skin and subcutaneous fat, with lesser involvement of the mus-
cles of mastication (Fig 2A). Skeletal findings were most pronounced in the mandible (Fig 2B). Enophthalmos of the left globe was present with relative paucity of retrobulbar fat (Fig 2C). No intracranial imaging findings were observed at the time of presentation.

Case 3. A 24-year-old man with progressive hemiatrophy of the left lower face and left facial nerve dysfunction during the past 3 years presented for imaging evaluation of symptoms. MR imaging demonstrated near-absence of subcutaneous fat involving the inframaxillary face extending inferiorly to the visualized portions of the suprathyroid neck (Fig 3A). Only minimal asymmetry was noted involving the muscles and skeletal structures (Fig 3B). Involved tissues demonstrated normal T1 and T2 signal without pathologic contrast enhancement (Fig 3C). Limited evaluation of intracranial content revealed no abnormality. Diagnosis of PRS was made, given imaging findings and compatible clinical history and examination.

Case 4. An 11-year-old girl with a known diagnosis of PRS of unknown duration was experiencing increasing dermal sensitivity and tightness of the left hemiface and occasional pain along the left temple for the past year. CT revealed soft-tissue atrophy and osseous asymmetry most prominently in the left maxilla (Fig 4A), with extension to the left frontal scalp superiorly and left mandible inferiorly. There was deviation of the nose toward the side of atrophy (Fig 4B). The maxillary sinus, ethmoidal air cells, mandible, and osseous orbit were relatively smaller on the left (Fig 4C). No dedicated intracranial imaging was performed.

Case 5. A 30-year-old woman with known PRS presented with weakness and recurrent headaches and underwent MR imaging of the brain. Subtle thinning of the subcutaneous skin and fat was seen along the right scalp (Fig 5A). On FLAIR and T2-weighted images, increased signal was present in the ipsilateral subinsular, temporal, and occipital lobe white matter and in the cerebral peduncle (Fig 5B, -C). Susceptibility-weighted imaging demonstrated innumerable punctate foci throughout the right temporal, frontal, and occipital lobes and in the cerebellum (Fig 5D). The left hemisphere was normal.

FIG 1. Axial CT images of a 44-year-old woman with PRS obtained for cosmetic surgical planning. A, There is prominent hemiatrophy of the skin, subcutaneous fat, and masseter muscle. B, The right maxillary sinus is considerably smaller than the left. C, Marked right enophthalmos is present.

FIG 2. A 14-year-old boy who presented with progressive atrophy of the left face. A, T1-weighted image shows striking paucity of fat in the left face. B, 3D reconstruction of a CT series demonstrates left facial hemiatrophy, particularly involving the mandible. C, T1-weighted image at the level of the orbits reveals left enophthalmos.
Case 6. A 35-year-old woman with known left PRS presented with headache and weakness. MR imaging demonstrated increased T2 signal in the left hemispheric white matter. Involvement was geographic, with regional areas of confluent periven-tricular, deep, and subcortical white matter hyperintensity (Fig 6). The right hemisphere demonstrated only nonspecific punctate foci of white matter hyperintensity.

Case 7. A 68-year-old man with known PRS presented with clinical acute right middle cerebral artery occlusion. Noncontrast head CT demonstrated marked thinning of the left frontal scalp similar to an en coup de sabre deformity (Fig 7A). MR imaging demonstrated findings consistent with right MCA acute infarction (not shown). The left hemisphere was markedly abnormal with extensive white matter signal abnormality (Fig 7B). Asymmetric left ventriculomegaly and a relatively small ipsilateral subcortical size (Fig 4C) were present and suggested hydrocephalus, but their etiology remained unexplained.

DISCUSSION
Parry Romberg syndrome is a sporadic and rare condition that has been reported to be more common in females, without apparent geographic or ethnic predilection. Onset typically occurs during the first and second decades of life, resulting in an initially insidious but progressive hemiatrophy of the face during a span of 2–20 years, with a slight propensity for the left side. Seemingly without cause, the progression abruptly arrests and stabilizes, reaching a “burned-out” phase. Diagnosis of PRS mainly relies on the clinical history and examination and exclusion of other possibilities, supported by histopathologic and imaging studies.

Patients characteristically experience atrophy of the skin and subcutaneous tissues and may develop atrophy of the underlying muscular, cartilaginous, osseous, and glandular structures as the disease progresses. The affected area typically begins in the maxillary or periorbital region and may expand to involve the forehead, perioral region, teeth, jaw, and neck to varying de-
Involvement of the teeth may help in determining the age of onset in unclear cases, because the presence of relatively smaller teeth with short roots has been reported in PRS. Earlier onset and longer duration of PRS have been reported to relate to increased severity of the disease. Superficial changes such as skin discoloration and cicatricial alopecia may occur as well. PRS is typically limited to 1 side of the head and neck, but bilateral disease has been reported. Similar manifestations involving the trunk and extremities are even rarer.

Neurologic symptoms occur in 15%–20% of patients, with the most common being ipsilateral headaches, facial pain, and seizures, which may be refractory to treatment. Other reported neurologic symptoms include trigeminal neuritis, facial paresthesia, cranial nerve dysfunction, fixed focal neurologic defects, hemiparesis, and cognitive impairment. Ophthalmologic symptoms occur in 10%–35% of patients and usually involve the ipsilateral orbit. Enophthalmos is common due to atrophy of the retrobulbar fat; other potential orbital abnormalities include uveitis and retinal or optic nerve alterations.

Many theories have developed during the years in an attempt to explain this peculiar disease. However, no single theory has been satisfactory in fully characterizing and predicting PRS, and our understanding of the underlying
pathophysiology remains limited. Proposed etiologies have included trauma, genetic predisposition, infection, radiation exposure, embryonic developmental dysfunction, sympathetic cervical ganglion dysfunction, and metabolic and endocrinologic disturbances. Currently, the strongest laboratory and histologic evidence supports an inflammatory autoimmune disorder with or without associated vasculopathy. Clinically, evidence suggests that a striking pattern of innumerable foci of susceptibility in the ipsilateral brain—these may represent microhemorrhages or foci of mineralization. In another patient, unexplained ventriculomegaly and associated small sulci could represent underlying hydrocephalus or may reflect an as-yet-undocumented intracranial manifestation of PRS.

PRS shares many clinical features, such as age of onset, associated neurologic symptoms, and cutaneous presentations with en coup de sabre, a rare variant of linear scleroderma that occurs along the anterior aspect of the scalp. These 2 entities are commonly discussed in conjunction due to their overlapping presentations and lack of distinguishing clinical features. To make differentiation even more complicated, conversion of en coup de sabre to PRS has been reported, and 30%–40% of patients with PRS have changes typical of scleroderma localized outside the face. This has led many authors to consider these 2 entities variants that share a common underlying pathogenesis related to linear scleroderma. Other authors, however, argue against this claim because deeper head and neck structures, such as the oral or pharyngeal musculature, are not commonly involved in en coup de sabre, and there are typically elevated titers of autoantibodies in en coup de sabre but not in PRS. On the other hand, PRS demonstrates little-to-no sclerosis of the cutaneous and subcutaneous structures, variable perivascular chronic inflammation, and preservation of dermal elastic tissue on histopathologic examinations, differentiating it from the spectrum of scleroderma diseases.

PRS also shares features with Rasmussen encephalitis, an autoimmune/inflammatory disorder affecting 1 side of the brain, resulting in refractory epilepsy and progressive hemiplegia and typically affecting children in the first decade of life. On imaging, Rasmussen encephalitis is characterized by predominately unilateral hemispheric areas of hyperintense T2 signal with eventual development of unilateral ce-

FIG 7. A 68-year-old man with PRS who presented with acute right MCA infarct. A, Axial CT image demonstrates striking focal loss of scalp thickness in the left frontal region. B, Axial FLAIR images demonstrate confluent signal abnormality in the high left frontal and parietal white matter. C, Etiology of left ventriculomegaly and sulcal asymmetry was uncertain.
trophic (Barraque-Simons syndrome) may have manifestations.

infectious, vitamin D3 analogues, and penicillamine are also treatment options that have demonstrated variable responses. Antimalarials, antibacterials, corticosteroids, immunomodulators, and plasmapheresis, have been used with varying degrees of success. Antimalarials, antibiotics, and plasmapheresis have been attempted as well due to concerns for cost and patient radiation exposure.23,24

Volumetric calculations can be easily achieved with 3D reconstructions of CT data acquired periprocedurally and on serial follow-up examinations, but alternative methods are being investigated as well due to concerns for cost and patient radiation exposure.23,24

In summary, PRS is a rare, self-limiting, and slowly progressive hemiatrophy of the face that typically affects the skin and subcutaneous tissues and may affect deeper tissues such as the musculature, cartilage, and osseous structures. Neurologic and ophthalmologic symptoms are common, but underlying pathophysiology remains uncertain. Intracranial involvement is best evaluated with MR imaging and can range widely, with the most common findings being parenchymal calcifications, white matter abnormalities, and brain atrophy. In addition to evaluating the extent of disease, radiologic assessments may also facilitate the exclusion of other differential considerations, help monitor disease progression, and evaluate posttreatment responses.

REFERENCES