MEDICAL GRAND ROUNDS

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The Problem of Osteoarthritis

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"Arthritis is a mean disease. It will not turn you blue (Though, if it did, the doctor's task were easier, 'tis true). It doesn't kill you suddenly - you're apt to live for years. It doesn't fill your family with sympathetic tears. The Early Briton suffered on his Early British cliff And told the local druids that his joints were feeling stiff."

(From a poem by Stephen Vincent Benet who suffered from arthritis.)

I. The Differential Diagnosis of Osteoarthritis.

<u>Case #1</u>: This 65-year-old WM was admitted the fibrillation, mitral and aortic murmurs and gave a history of rheumatic fever in 1917. A diagnosis of rheumatic heart disease was made. In 1949 he first complained of pain in his fingers and hips. His told him he had rheumatoid arthritis and started him on cortisone in 1952. He obtained some relief and continued thereafter on the appropriate fashionable steroid although troublesome edema was induced by some of the tablets. At the time of admission, he was taking 10 mg. prednisone daily. Examination showed swelling of the DIP's and PIP's and marked limitation of the left hip movements. X-rays revealed classical changes of osteoarthritis and rheumatoid factor tests were negative.

Case #2: This 52-year-old WF presented on 759 with a one year history of pain in the hands, shoulders, knees, ankles and feet plus a chart of her family tree for six generations showing 9 female relatives with arthritis. The PIP's and DIP's were swollen, right shoulder movements were limited, and a small effusion was noted in the left knee. The latex and sensitized sheep cell tests were negative, ESR 20, uric acid 5 mg%. She was at first considered to have rheumatoid arthritis. However, later examination showed Heberden's and Bouchard's nodes, and the x-rays osteoarthritic changes in the DIP's, PIP's and right shoulder. At present she is doing well and has had no further joint involvement.

II. Arthritis as a Diagnostic Clue

case #3: This WF was first seen here in y 1962 at the age of 44 c/o visual difficulties. She had the following history: She was in good health until 1949. Pain both knees and many PIP joints. Noticed increase in finger size and had wedding ring cut off. 1952. Menorrhagia and gyn. op. ?type. 1953. Weight loss, polyphagia and polydyspia. Diabetes diagnosed and insulin therapy commenced. About this time, joint pains worsened and had to increase shoe size $6 \rightarrow 9$. She began to have severe headaches. Acromegaly diagnosed and patient given pituitary irradiation. 1954. She remained asymptomatic until 1962 when, during the second states, she noticed pain in the hands and ribs. She had a further course of irradiation with improvement in the joint symptoms but visual difficulties and fatigue continued. On admission to on 62, she had obvious acromegaly. X-rays revealed a very enlarged sella turcica, thickening of the shafts of the phalanges of both hands, with tufting of the PIP's, tufting of the distal portions of the metatarsals of both feet, and changes of degenerative joint disease in the thoracic spine, She has had lens extractions and remains relatively well.

III. ?Cervical Spondylosis

Case #4: The formation of the left arm. This 66-year-old CM was first seen 1957 with a "spastic gait." He had been well until 1953, when he developed pain and swelling in the left ankle, hips, and right thumb. Some time later, his neck was painful and stiff and he noticed weakness of the left arm. He was admitted to and given "shots." By 1956, he had marked difficulty in walking and by 1957 was using sticks. His first thorough work-up was in 1962. The BP was 180/110 but the rest of the chest, heart and abdominal examination was WNL. The CNS findings could be summarized as a spastic tetraplegia, more marked on the left side with atrophy of the arm musculature. There was no real evidence of peripheral joint involvement. The following diagnoses were considered during the next few months: Multiple sclerosis, motor neurone disease, basilar artery insufficiency. An x-ray of the cervical spine was taken 1962, and showed marked changes. He was then admitted for evaluation.

Laboratory tests showed Hb 12.6, ESR 36 mm/hr., globulin 4 G%, ceph flocc 3+, latex (slide test) 2+, SCAT, Antinuclear test and serology neg. Extensive x-rays of the cervical spine showed disappearance of all disk spaces, osteoporosis of vertebral bodies, juxta-articular spurring esp. of C 6 and 7, and narrowing of many neural foramina, esp. upper cervical region. There was also an abnormal relationship between the odontoid and C 1, with marked anterior displacement of C 1, resulting in reduction of the caliber of the neural canal. The right atlanto-occipital joint was narrowed. Clinically, there was complete loss of the cervical curve. The remainder of the skeletal survey showed generalized demineralization with slight osteoarthritic changes in the lumbar spine and hips. The hands showed cystic changes with subluxation of the proximal phalanges right thumb. The CSF was normal except that protein was 76 mg%. Myelogram was unsuccessful as the dye would not fill the cervical canal. A pneumoencephalogram showed filling of the ventricles. Electromyography confirmed a widespread neuropathic lesion with frank denervation in C-5 region. He was discharged without a definite diagnosis, and was last seen $\frac{1}{100}/63$ when there was little objective change. He was, however, subjectively improved.

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Malformations, malalignments of cravitatic, congenital or neoplastic origin
Pulmonary cateoarthropathy Herophilia
Gout Neuropathies (e.g. tabes formalis and syringuryelia)

TABLE 2.

Specific Forms of Osteouchritis

 Roberden's nodes ("What are those little hard knobs, shout the size of a pea, which are frequently seen upon the fingers, a little below the top, sear the joint." William Heberden, 1802)

Primary generalized osteoarthritis (F.C.C.)

. Knees

. Hips

5. Spine

TABLE 1.

Classification

primary osteoarthritis

Causes: Degeneration ?Genetic ?Other

secondary osteoarthritis

Trauma (inc. contusions, intra- or juxta-articular Causes: fractures or dislocations) Obesity Malformations, malalignments of traumatic, congenital or neoplastic origin Infections and inflammatory joint diseases Osteochondritis and aseptic necrosis of epiphyseal bone Pulmonary osteoarthropathy Hemophilia Hemochromatosis Ochronosis Acromegaly Muscle abrophy Gout Neuropathies (e.g. tabes dorsalis and syringomyelia)

TABLE 2.

Specific Forms of Osteoarthritis

- Heberden's nodes ("What are those little hard knobs, about the size of a pea, which are frequently seen upon the fingers, a little below the top, near the joint." William Heberden, 1802)
- 2. Primary generalized osteoarthritis (P.G.O.)
- 3. Knees
- 4. Hips of disease
- 5. Spine

Usually good as regards function unless hips involved

Cummonest in Ruses

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Differential Diagnosis o	f Rheumatoid Arthritis	vs. Osteoarthritis*
Age of onset	Usually 20-45 years	Usually over 40 years
Hands	Proximal inter- phalangeal and meta- carpophalangeal	Distal and proximal interphalangeal joints
	joints	First carpo-metacarpal joints
Other joints	Any joint may be affected	Knees, hips, sacroiliac, lumbar and lower cervical
Number joints	Many	Few (except in P.G.O.)
Type onset	Usually insidious; may be acute	Insidious
Constitutional symptoms	Yes	Usually none
Type joint swelling	Usually fusiform or spindle shaped	Usually irregular or knobby
Muscle atrophy	Usually present and may be marked	Infrequent
Taink a Officer i an	The survey of	
Joint errusion	Frequent	Inirequent
		Commonest in knees
Subcutaneous nodules	15%	Absent
ESR 1. Motor neurone di	Usually elevated	Usually normal
2. Nultiple scleros 3. Syringomyelia 4. Syphilitic disea		Occasionally slightly elevated
Rheumatoid factor (by standard tests)	80% of cases	Negative, except when diseases associated
		tors present
Progress of disease	Marked tendency to progression	Slow progress
Prognosis	Variable	Usually good as regards function unless hips

-5-

Rheumatoid Arthritis vs. Osteoarthritis

X-ray Findings Early

Periarticular soft tissue swelling

Rarefaction of trabeculated ends of bones

Later

Progressive narrowing joint space

Cortical erosions with punched out areas of bone at articular margins

Subluxations and dislocations

Ankylosis

Narrowing joint space

Formation bony spicules and osteophytes at margins of joints

Cysts at bone ends

Condensation of subchondral bone

Deformities from pressure and trac-

No ankylosis

*Adapted from "Arthritis," ed. J. L. Hollander.

Denoted of blood vessels and 1 TABLE 4.

Differential Diagnosis of Cervical Spondylosis

A. Cord Compression

- 1. Motor neurone disease
 - 2. Multiple sclerosis
 - 3. Syringomyelia
 - 4. Syphilitic disease in its various forms

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- 5. Sub-acute combined degeneration
- 6. Spinal cord tumors

B. Root Compression

- 1. Primary and metastatic tumors
- 2. Syringomyelia
- 3. Median nerve compression
 - 4. Peri-capsulitis of shoulder joint
- 5. Syndromes of the thoracic inlet and cervico-axillary canal
- 6. Shoulder-girdle neuritis
 - 7. Angina, coronary occlusion

TABLE 5.

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17

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Presenting Symptom or Syndrome in 100 Patients with Cervical Spondylosis (Brain, ref. 29)

Brachial radiculitis Headache Vertigo Myelopathy Pain in neck Vertebro-basilar ischemia Attacks of unconsciousness Drop attacks Acute lesion of spinal cord

TABLE 6.

toclassified phagoayteArticular Cartilage

Hyaline type in synovial joints. Average thickness 1 to 7 mm.

Aneural.

Devoid of blood vessels and lymphatics except in deepest layer adjacent to bone. Osteophyte formation occurs here.

Has no perichondrium, not covered by synovial membrane. Never ossifies.

Consists of cells, the chondrocytes, which lie in small cavities distributed in intercellular matrix.

Chondrocytes contain glycogen and lipids.

Mitotic cell division does not occur in adult cartilage.

Matrix consists of collagenous fibers in homogeneous ground substance.

Matrix contains high proportion of the mucopolysaccharide, chondroitin sulphuric acid with a polypeptide.

The chondroitin sulphate consists of a disaccharide unit containing glucoronic acid and glucosamine. Only types A and C chondroitin sulphate in cartilage. Probable molecular weight 260,000. Mucopolysaccharide imparts toughness and resilience to the tissue.

Ratio of chondromucoprotein to collagen highest in those joints which withstand the most pressure.

Metabolic activity of cartilage decreases with aging. In osteoarthritis, changes occur first in the superficial layer of cartilage.

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TF	7DT	12	- (•

The Compositio	n of	Nor	mal	Sync	vial	Flui	<u>d</u> *			
					25	Rang	е			Average
Amount in knee, ml	58	2 1	g.	i zist	0	.13-	3.5	2		1.1
Relative viscosity at 25°C					5	.7-1	160			235
pH					7	.2-7	•4			7.4
Leukocyte count, per cu mm						13-1	80			63
Differential, per cent										
Polymorphonuclears						0-2	5			6.5
Lymphocytes						0-7	8			24.6
Monocytes						0-7	1			47.9
Clasamatocytes		. 8				0-2	6			10.1
Unclassified phagocytes						0-2	1			4.9
Synovial lining cells						0-1	2			4.3
Total solids, gm per 100 gm					2	•4-4	.83			3.41
Total albumin and globulin, gm	per	100	ml		1.0	07-2	.13			1.72
Albumin, gm per 100 ml										1.02
Globulin, gm per 100 ml						23				0.05
Mucin, nitrogen, gm per 100 ml					0.00	68-0	.135			0.104
Mucin glucosamine, gm per 100 m	ml				0.0	12-0	.132			0.074
Fibrinogen						0				
Glucose				Ap	prox.	the	same	as	in j	plasma
Nonprotein nitrogen					11	11	н	11	1	11
Uric acid					u Ö	11	11	11	II S	"
Electrolytes				Ap	prox.	the	same	as	pla	sma

Approx. the same as plasma dialysate

*Modified from M.W. Ropes and W. Bauer: Synovial Fluid Changes in Joint Disease, Harvard University Press, Cambridge, Mass., 1953.

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		R, M Marici	B F Rhe or	060	l adel nhia	Pasan. Ph	a " R.A. Je	"Synovianalys"	*Adanted fro
+	0	0	ບ. ອ	Some- times bloody	25,000 to 100,000 85% polys	Poor	Low	Grayish, creamy bloody; cloudy	Septic Arthritis
+	0	ed 1 ta. ' O	ប ů	1961 • • • • • •	20,000 + 60% polys	Poor	Low	Yellow; cloudy	Tuberculous Arthritis
0	0	o 0	3.7	0	10,000 <u>+</u> 50% pol y s	Good	Decreased	Yellow; sl. cloudy	Rheumatic Fever
0	0	o tors O	4.7	Occa- sional	15,000 <u>+</u> 70% pol y s	Poor	Low	Yellow to green; cloudy	Rheumatoid Arthritis
0	0	in ort +	0.1	olo	13,000 + 70% pol y s	Poor	Decreased	Yellow to milky; cloudy to turbid	Gout
0	‡	0	ω •0	0	700 <u>+</u>	Good	High	Yellow to straw; clear	Osteo- arthritis
0	0 or +	0 0	4.2	Few to grossly bloody	1,500 +	Good	Moderately High	Yellow to bloody; cloudy	Traumatic
0	0	0	μ. υ	0	100 25% polys	Good	High	Straw; clear	Norma.1
Bac- ceria	Cartilage Fibrils t	Urate Crys- tals	Protein (gm%)	RBC	wBC	Mucin Clot Ropes Tes	Viscosity (Appearance	
				T FILLA	s or synovia	TSOUGETC	TTTTELEULTAT		

TABLE 8

Differential Diagnosis ארייים בייאאייי

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