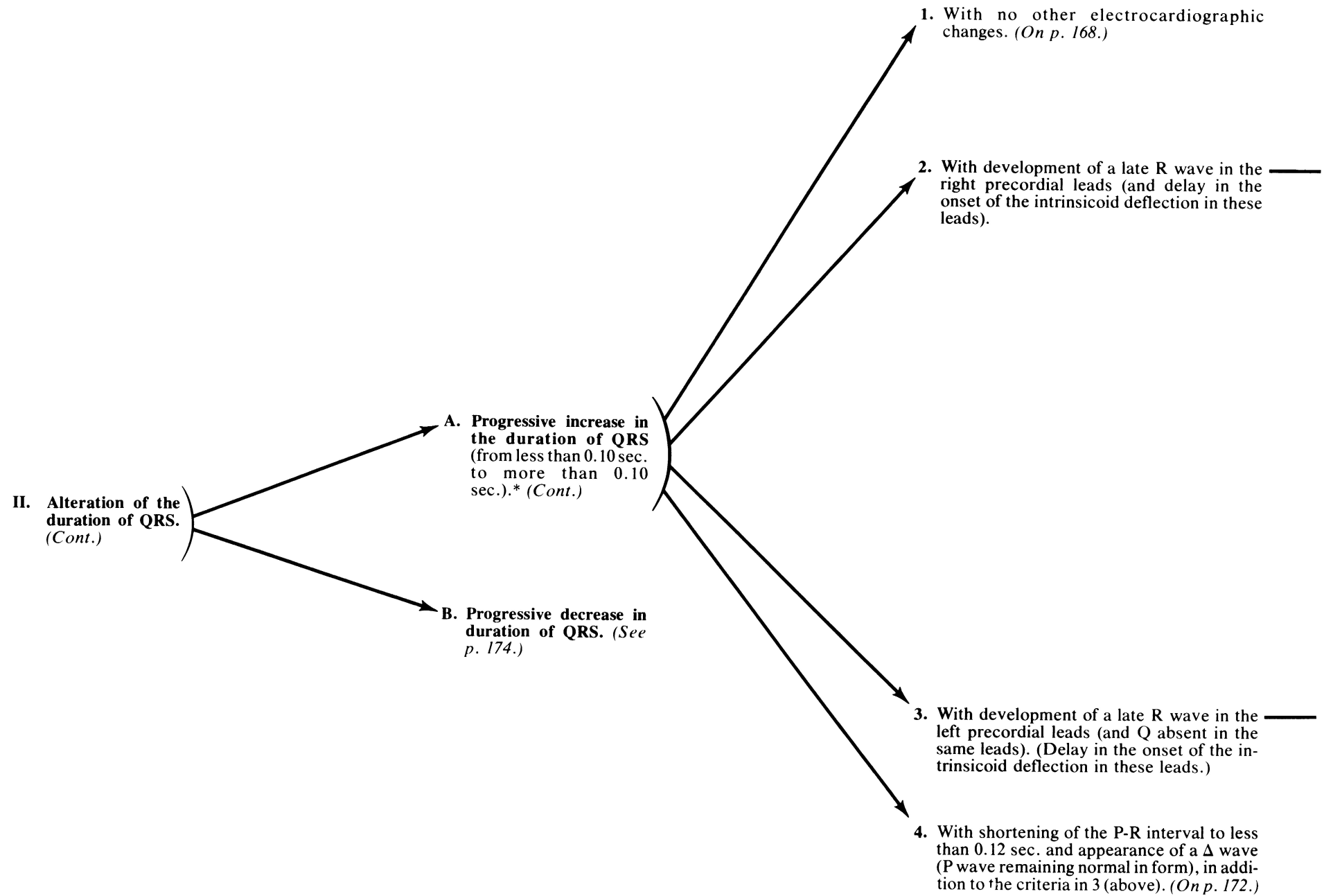


ACUTE SERIAL CHANGES:
Primarily limited to the QRS Complex (Cont.)



*Prolongation of QRS frequently leads to S-T and T changes.

†Complete or incomplete bundle branch block may appear during a run of supraventricular tachycardia or during an irregular rhythm when it exceeds a critical rate. This may be due to fatigue of either of the bundles.

Complete or incomplete **right bundle branch block**.

May be due to:

1. Myocardial infarction
2. Tachycardia†
3. Drug toxicity (digitalis, quinidine, procaine amide, etc.)
4. Acute cor pulmonale
5. Following right ventriculotomy
6. Injury or disease of the right bundle

FIG. Ser10A
control

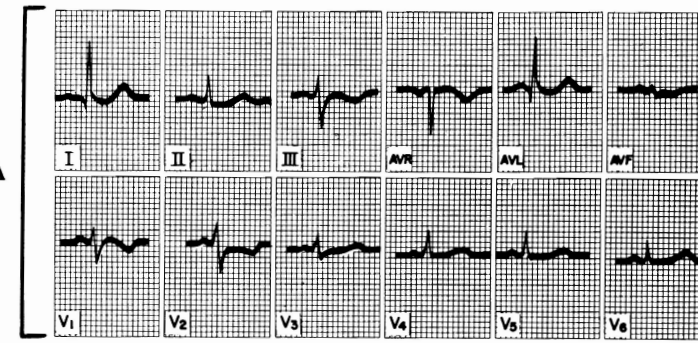
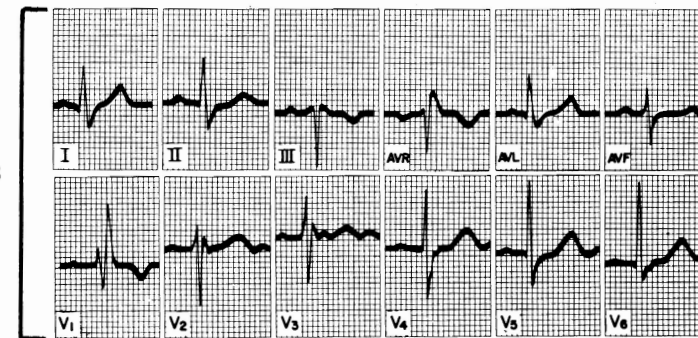


FIG. Ser10B



Complete or incomplete **left bundle branch block**.

Most often due to:

1. Myocardial infarction
2. Secondary to drug toxicity
3. Tachycardia†
4. Injury or disease of the left bundle
5. Electrolyte abnormalities

FIG. Ser11A
control

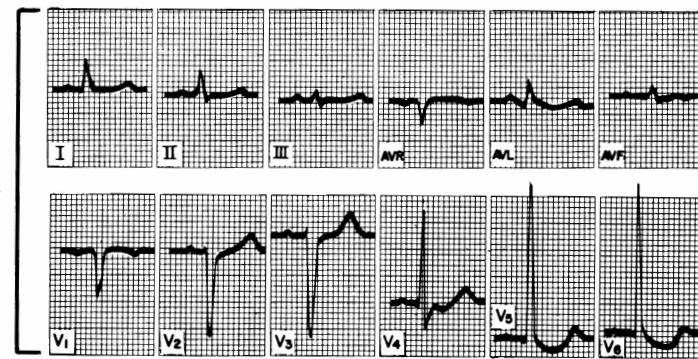
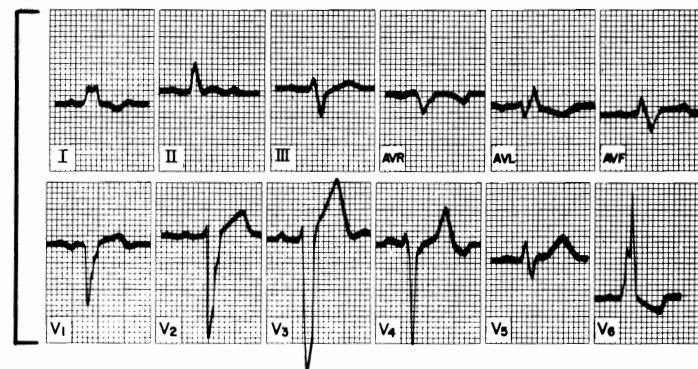
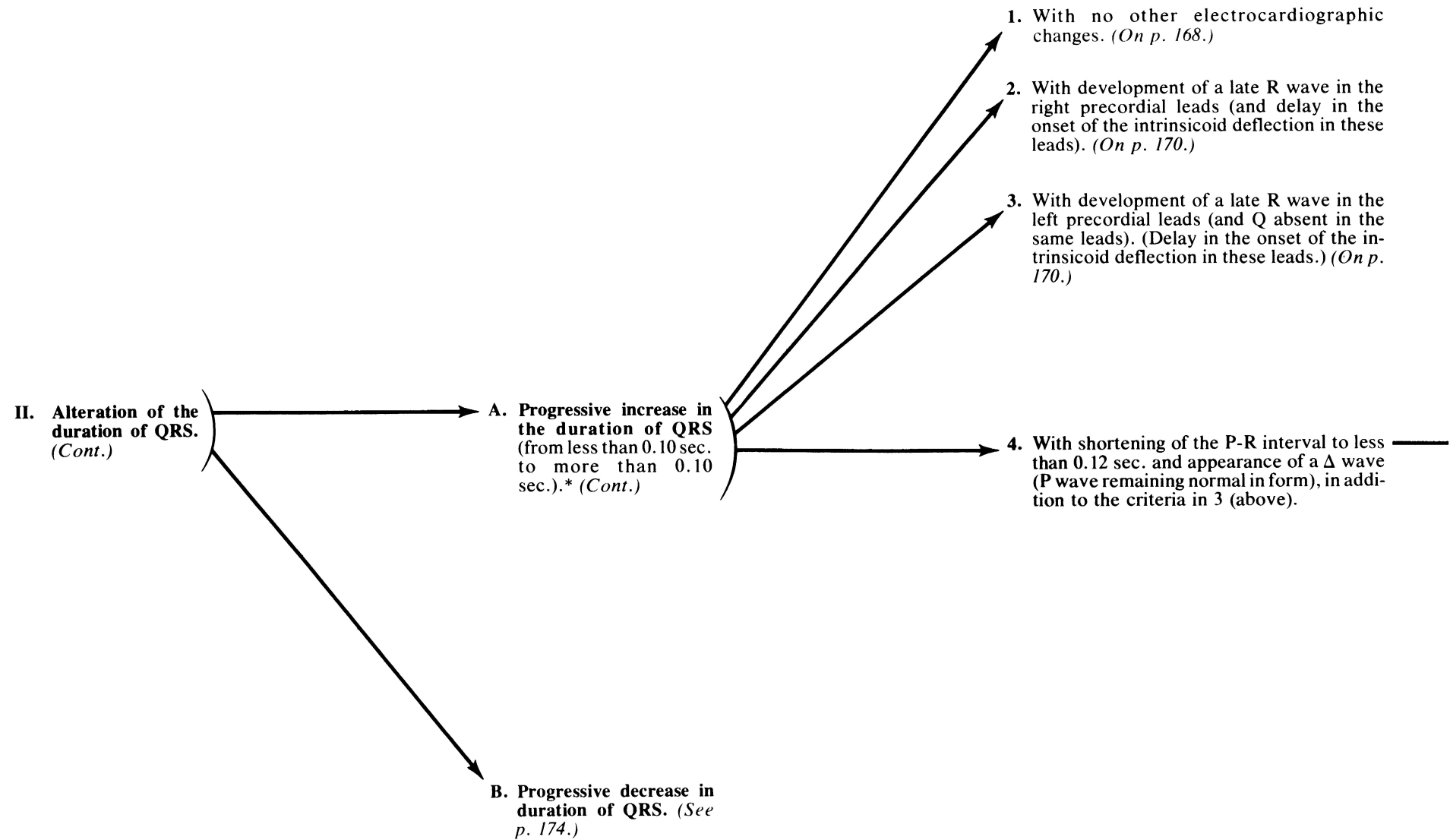


FIG. Ser11B



ACUTE SERIAL CHANGES:
Primarily limited to the QRS Complex (Cont.)



→ *Development of Wolff-Parkinson-White syndrome. (Shift from normal conduction to pre-excitation pathway.)*

FIG. Ser12A

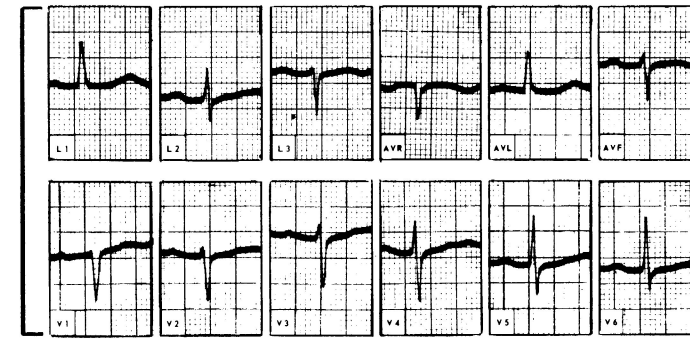
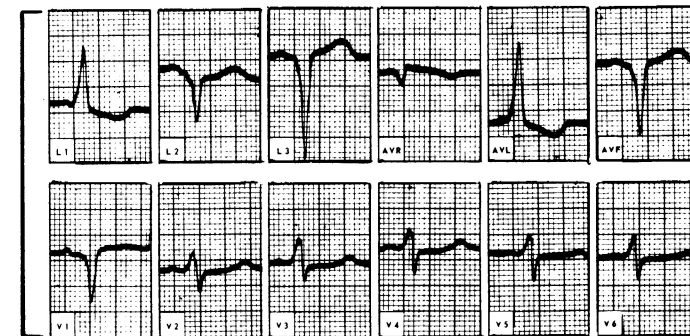
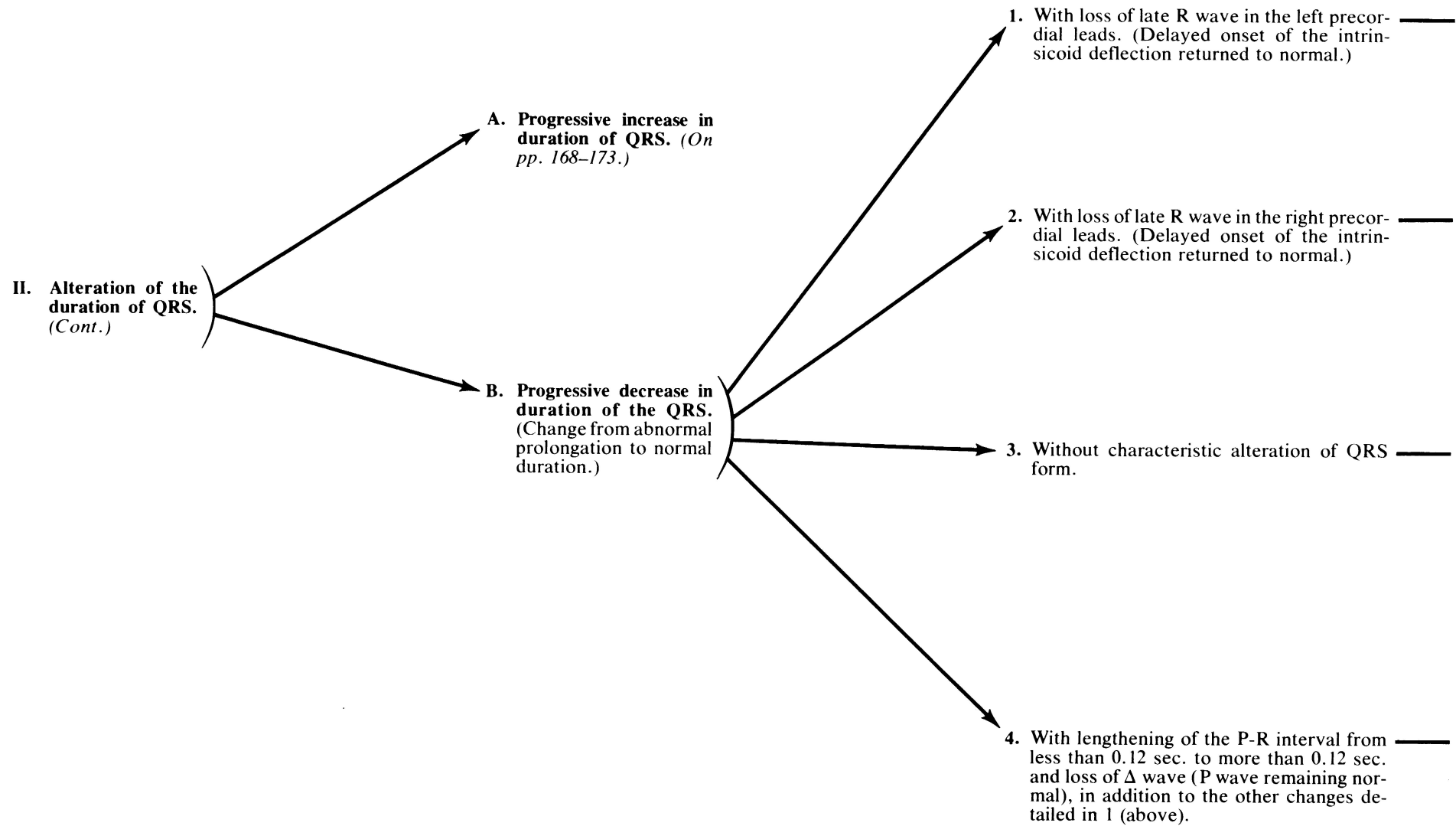


FIG. Ser12B



ACUTE SERIAL CHANGES:
Primarily limited to the QRS Complex (Cont.)



*Complete or incomplete bundle branch block may appear during a run of supraventricular tachycardia. This may be associated with fatigue of the bundles. See Chapter 2.

→ Revision of **left bundle branch block** to normal conduction. Occasionally seen during recovery from an **acute myocardial infarction**. More commonly seen following cessation of toxic action of drugs or the termination of a **tachycardia*** (Fig. Ser 11 in reverse-B to A).

→ Revision of **right bundle branch block** to normal conduction. Most commonly seen following the cessation of toxic action of drugs or the cessation of tachycardia* or during recovery from an acute **cor pulmonale**. Occasionally seen during recovery from acute myocardial infarction (Fig. Ser 10 in reverse-B to A).

→ Reversion of **intraventricular conduction defect** to normal. Seen with cessation of drug therapy, **correction of electrolyte imbalance**, or termination of tachycardia. On occasion seen during the healing of a **myocardial infarct**.

→ Change in atrioventricular conduction in **Wolff-Parkinson-White syndrome** from pre-excitation to normal.

FIG. Ser13A
control

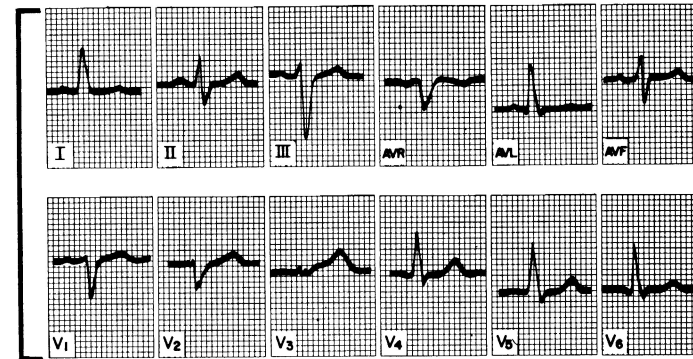


FIG. Ser13B

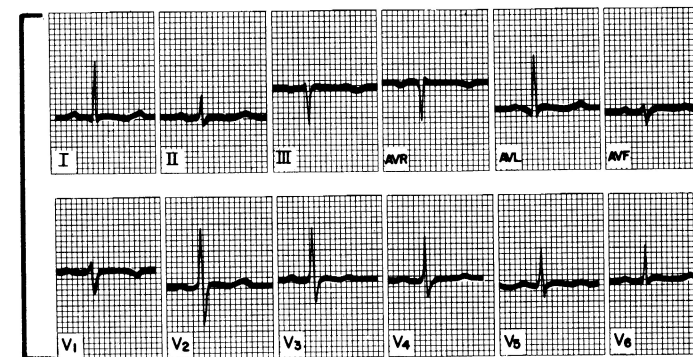


FIG. Ser14A

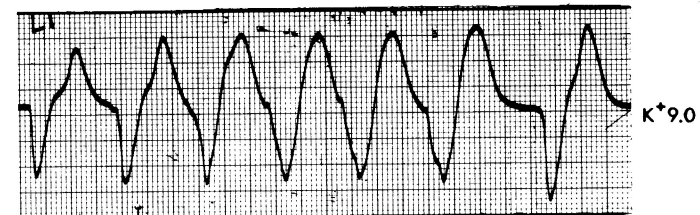


FIG. Ser14B

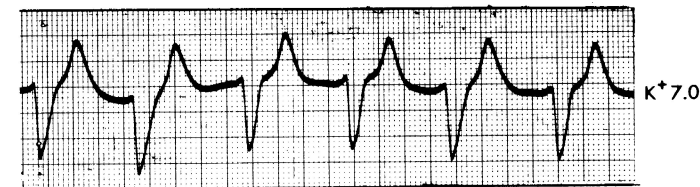
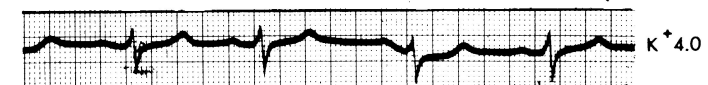


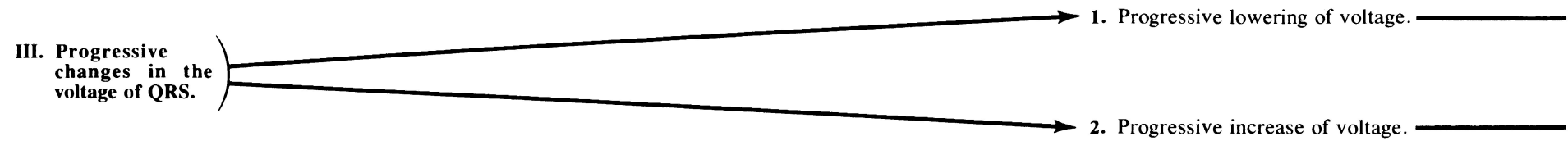
FIG. Ser14C



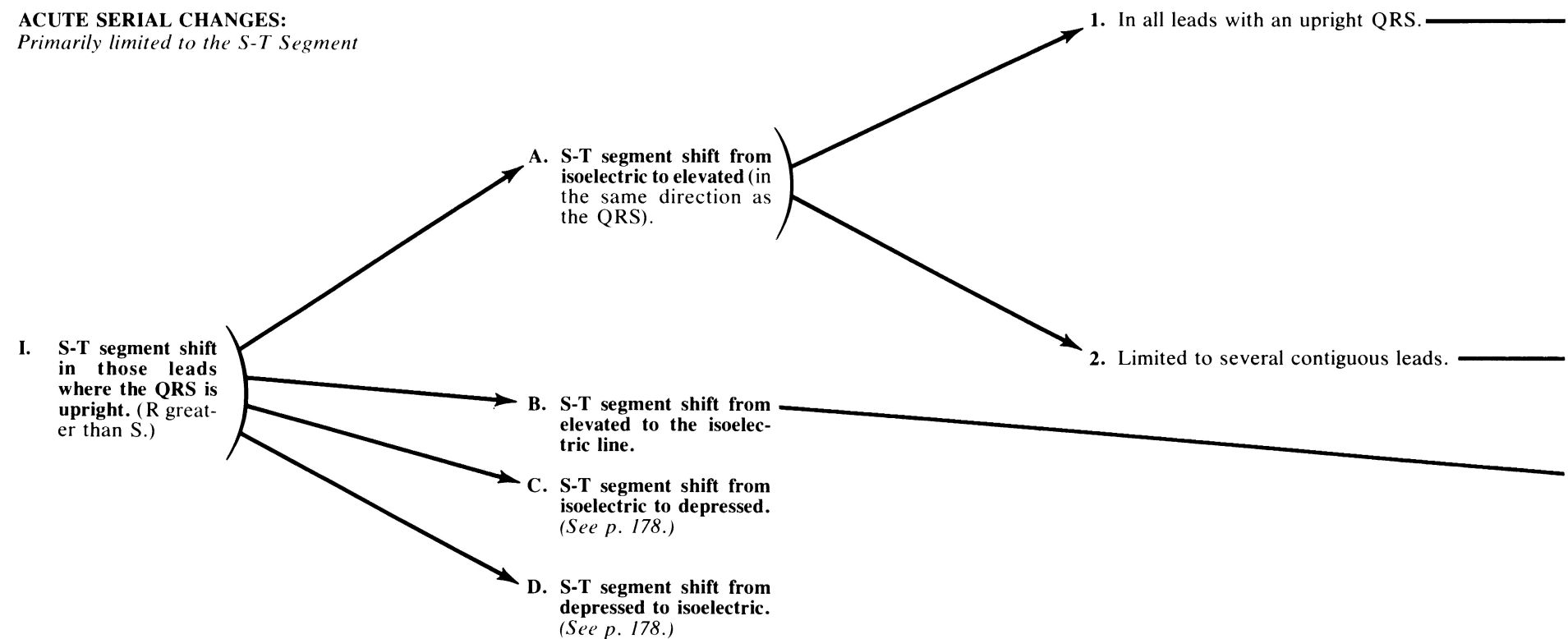
FIG. Ser14D



ACUTE SERIAL CHANGES:
Primarily limited to the QRS Complex (Cont.)



ACUTE SERIAL CHANGES:
Primarily limited to the S-T Segment



→ Pericardial effusion; massive pleural effusion; occasionally seen during the course of an acute myocardial infarction, but other changes are usually more characteristic. **Improper standardization of the ECG machine.**

→ With recovery from myocardial infarction; with absorption or removal of pericardial effusion; in rapid correction of myxedema. **Improper standardization of the ECG machine.**

FIG. Ser15A
control

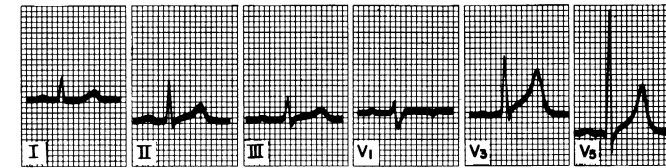
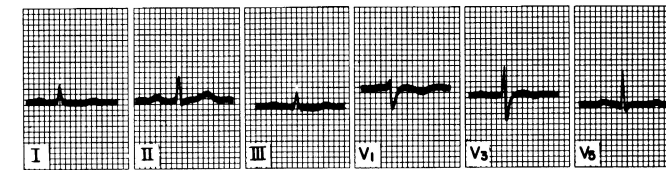
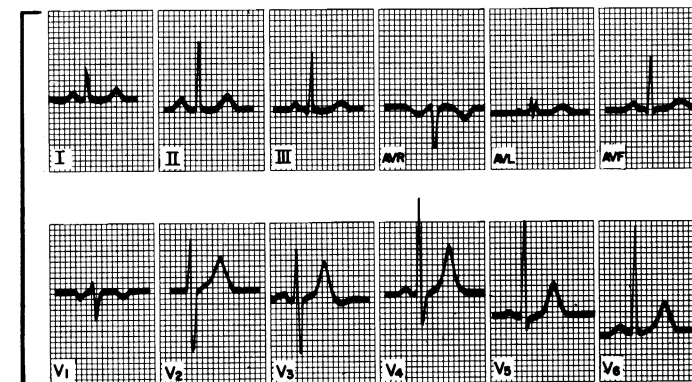


FIG. Ser15B



→ Acute pericarditis.

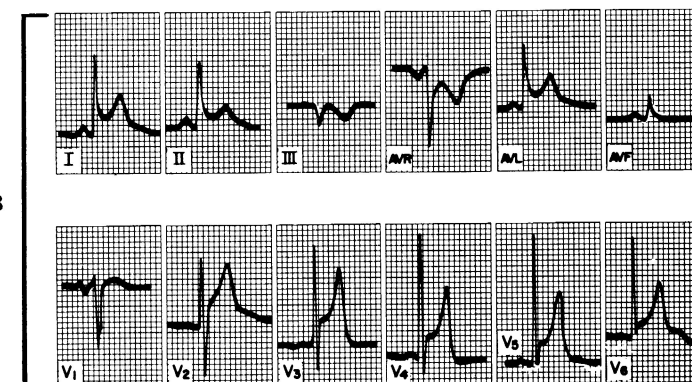
FIG. Ser16A
control



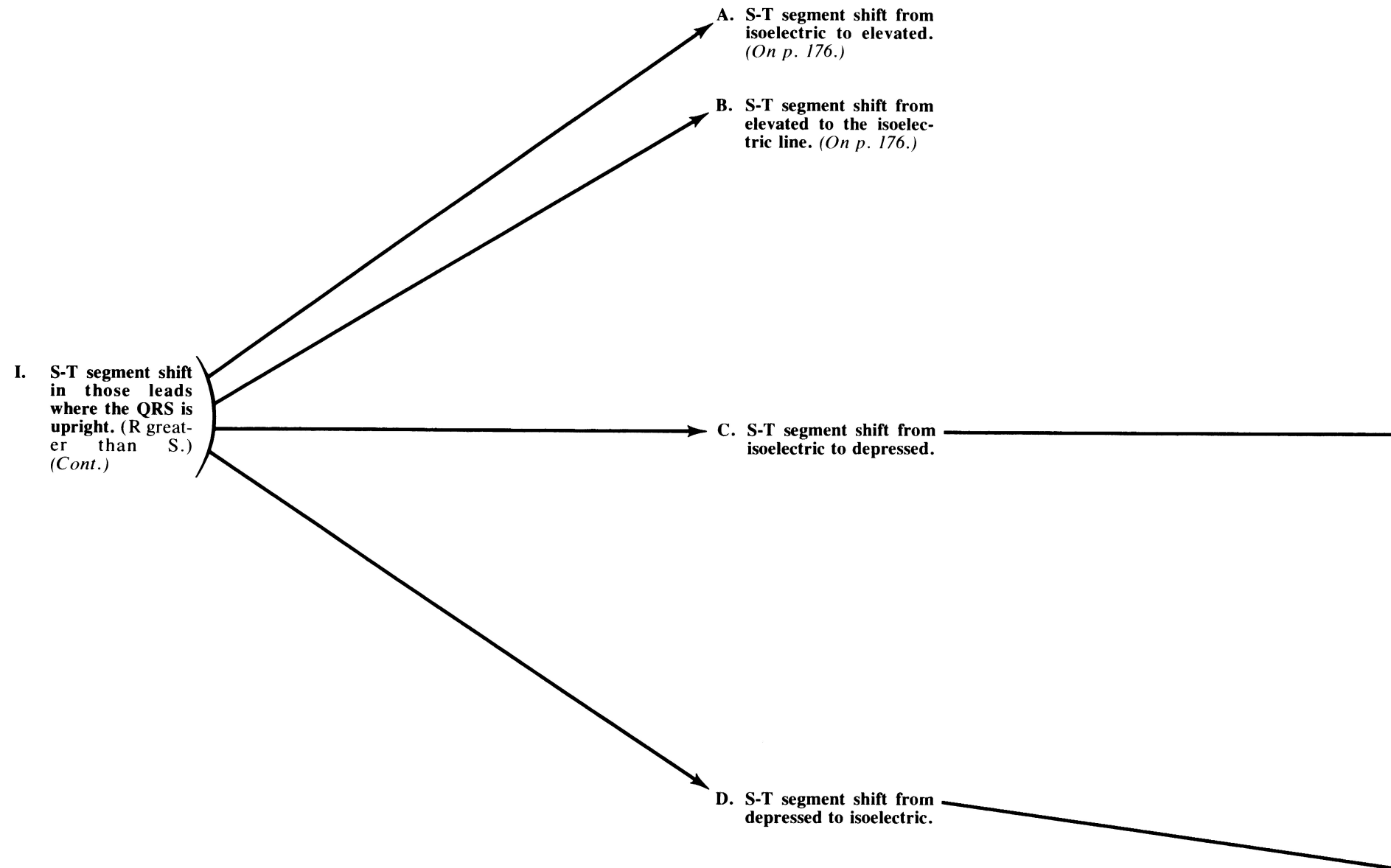
→ Possible early phase in the development of the electrocardiographic pattern of **myocardial infarction**. **Variant form of angina pectoris**. Possible localized **acute pericarditis**.

→ Serial change secondary to previously elevated S-T segment. Healing stage of pericarditis, termination of angina attack, etc.

FIG. Ser16B



ACUTE SERIAL CHANGES:
Primarily limited to the S-T Segment (Cont.)



*The differential diagnosis of the patterns of myocardial ischemia depends to a great extent on the clinical findings, the associated electrocardiographic changes, and the duration of these abnormalities.

A nonspecific change which may be found in any of the following clinical conditions:

1. Digitalis effect.
(During the administration of digitalis any diagnosis of other conditions as a cause of the depressed S-T, with or without T inversion, should be made with caution.)

FIG. Ser17A
control

FIG. Ser17B

2. Hypokalemia.

FIG. Ser18A
control

FIG. Ser18B

3. Myocardial ischemia* secondary to:
 - a. Angina pectoris (S-T segment changes persist for a few minutes to a half hour)*
 - b. Coronary insufficiency (S-T changes usually persist for several hours to one or two days)*
 - c. Subendocardial infarction (S-T changes usually persist several days or more)*

FIG. Ser19A
control

FIG. Ser19B

4. Tachycardia.
5. Acute cor pulmonale. (See Fig. Ser2.)
6. Drugs: quinidine, procaine amide, etc.
7. Defect in ECG machine (overdamped) or monitor tracing, compared with standard tracing.
8. Nonspecific; anemia, shock, cerebrovascular accidents, etc.

FIG. Ser20A

FIG. Ser20B

FIG. Ser21A

FIG. Ser21B

A serial change secondary to a previously depressed S-T segment. Recovery from, correction, or cessation of any of the above causes is followed by the return of the S-T segment to normal.

