In adults a tachycardia is any heart rate greater than 100 beats per minute. Supraventricular tachycardias may be divided into two distinct groups depending on whether they arise from the atria or the atrioventricular junction. This article will consider those arising from the atria: sinus tachycardia, atrial fibrillation, atrial flutter, and atrial tachycardia. Tachycardias arising from re-entry circuits in the atrioventricular junction will be considered in the next article in the series.

Clinical relevance

The clinical importance of a tachycardia in an individual patient is related to the ventricular rate, the presence of any underlying heart disease, and the integrity of cardiovascular reflexes. Coronary blood flow occurs during diastole, and as the heart rate increases diastole shortens. In the presence of coronary atherosclerosis, blood flow may become critical and anginal-type chest pain may result. Similar chest pain, which is not related to myocardial ischaemia, may also occur. Reduced cardiac performance produces symptoms of faintness or syncope and leads to increased sympathetic stimulation, which may increase the heart rate further.

As a general rule the faster the ventricular rate, the more likely the presence of symptoms—for example, chest pain, faintness, and breathlessness. Urgent treatment is needed for severely symptomatic patients with a narrow complex tachycardia.

Electrocardiographic features

Differentiation between different types of supraventricular tachycardia may be difficult, particularly when ventricular rates exceed 150 beats/min.

Knowledge of the electrophysiology of these arrhythmias will assist correct identification. Evaluation of atrial activity on the electrocardiogram is crucial in this process. Analysis of the ventricular rate and rhythm may also be helpful, although this rate will depend on the degree of atrioventricular block. Increasing atrioventricular block by manoeuvres such as carotid sinus massage or administration of intravenous adenosine may be of diagnostic value as slowing the ventricular rate allows more accurate visualisation of atrial activity. Such manoeuvres will not usually stop the tachycardia, however, unless it is due to re-entry involving the atrioventricular node.

**Sinus tachycardia**

Sinus tachycardia is usually a physiological response but may be precipitated by sympathomimetic drugs or endocrine disturbances.

The rate rarely exceeds 200 beats/min in adults. The rate increases gradually and may show beat to beat variation. Each P wave is followed by a QRS complex. P wave morphology and axis are normal, although the height of the P wave may increase with the heart rate and the PR interval will shorten. With a fast tachycardia the P wave may become lost in the preceding T wave.
Recognition of the underlying cause usually makes diagnosis of sinus tachycardia easy. A persistent tachycardia in the absence of an obvious underlying cause should prompt consideration of atrial flutter or atrial tachycardia.

Rarely the sinus tachycardia may be due to a re-entry phenomenon in the sinoatrial node. This is recognised by abrupt onset and termination, a very regular rate, and absence of an underlying physiological stimulus. The electrocardiographic characteristics are otherwise identical. The rate is usually 130-140 beats/min, and vagal manoeuvres may be successful in stopping the arrhythmia.

### Atrial fibrillation

This is the most common sustained arrhythmia. Overall prevalence is 1% to 1.5%, but prevalence increases with age, affecting about 10% of people aged over 70. Causes are varied, although many cases are idiopathic. Prognosis is related to the underlying cause; it is excellent when due to idiopathic atrial fibrillation and relatively poor when due to ischaemic cardiomyopathy.

Atrial fibrillation is caused by multiple re-entrant circuits or “wavelets” of activation sweeping around the atrial myocardium. These are often triggered by rapid firing foci. Atrial fibrillation is seen on the electrocardiogram as a wavy, irregular baseline made up of f (fibrillation) waves discharging at a frequency of 350 to 600 beats/min. The amplitude of these waves varies between leads but may be so coarse that they are mistaken for flutter waves.

Conduction of atrial impulses to the ventricles is variable and unpredictable. Only a few of the impulses transmit through the atrioventricular node to produce an irregular ventricular response. This combination of absent P waves, fine baseline f wave oscillations, and irregular ventricular complexes is characteristic of atrial fibrillation. The ventricular rate depends on the degree of atrioventricular conduction, and with normal conduction it varies between 100 and 180 beats/min. Slower rates suggest a higher degree of atrioventricular block or the patient may be taking medication such as digoxin.

Fast atrial fibrillation may be difficult to distinguish from other tachycardias. The RR interval remains irregular, however, and the overall rate often fluctuates. Mapping R waves against a piece of paper or with calipers usually confirms the diagnosis.

Atrial fibrillation may be paroxysmal, persistent, or permanent. It may be precipitated by an atrial extrasystole or result from degeneration of other supraventricular tachycardias, particularly atrial tachycardia and/or flutter.

### Atrial flutter

Atrial flutter is due to a re-entry circuit in the right atrium with secondary activation of the left atrium. This produces atrial contractions at a rate of about 300 beats/min—seen on the electrocardiogram as flutter (F) waves. These are broad and appear saw-toothed and are best seen in the inferior leads and in lead V1.

The ventricular rate depends on conduction through the atrioventricular node. Typically 2:1 block (atrial rate to

---

### Causes of sinus tachycardia

**Physiological**—Exertion, anxiety, pain

**Pathological**—Fever, anaemia, hypovolaemia, hypoxia

**Endocrine**—Thyrotoxicosis

**Pharmacological**—Adrenaline as a result of phaeochromocytoma; salbutamol; alcohol, caffeine

---

### Causes of atrial fibrillation

- Ischaemic heart disease
- Hypertensive heart disease
- Rheumatic heart disease
- Thyrotoxicosis
- Alcohol misuse (acute or chronic)
- Cardiomyopathy (dilated or hypertrophic)
- Sick sinus syndrome
- Post-cardiac surgery
- Chronic pulmonary disease
- Idiopathic (lone)
ventricular rate) occurs, giving a ventricular rate of 150 beats/min. Identification of a regular tachycardia with this rate should prompt the diagnosis of atrial flutter. The non-conducting flutter waves are often mistaken for or merged with T waves and become apparent only if the block is increased. Manoeuvres that induce transient atrioventricular block may allow identification of flutter waves.

Atrial flutter (rate 150 beats/min) with increasing block (flutter waves revealed after administration of adenosine)

The causes of atrial flutter are similar to those of atrial fibrillation, although idiopathic atrial flutter is uncommon. It may convert into atrial fibrillation over time or, after administration of drugs such as digoxin.

Atrial flutter with variable block

The ventricular rate depends on the degree of atrioventricular block, but when 1:1 conduction occurs a rapid ventricular response may result. Increasing the degree of block with carotid sinus massage or adenosine may aid the diagnosis.

Atrial tachycardia typically arises from an ectopic source in the atrial muscle and produces an atrial rate of 150-250 beats/min—slower than that of atrial flutter. The P waves may be abnormally shaped depending on the site of the ectopic pacemaker.

Atrial tachycardia with 2:1 block (note the inverted P waves)

Atrial tachycardia is initiated by an ectopic atrial focus (the P wave morphology therefore differs from that of sinus rhythm)

Types of atrial tachycardia
- Benign
- Incessant ectopic
- Multifocal
- Atrial tachycardia with block (digoxin toxicity)

The ventricular rate depends on the degree of atrioventricular block, but when 1:1 conduction occurs a rapid ventricular response may result. Increasing the degree of block with carotid sinus massage or adenosine may aid the diagnosis.

There are four commonly recognised types of atrial tachycardia. Benign atrial tachycardia is a common arrhythmia in elderly people. It is paroxysmal in nature, has an atrial rate of 80–140 beats/min and an abrupt onset and cessation, and is brief in duration.
Incessant ectopic atrial tachycardia is a rare chronic arrhythmia in children and young adults. The rate depends on the underlying sympathetic tone and is characteristically 100-160 beats/min. It can be difficult to distinguish from a sinus tachycardia. Diagnosis is important as it may lead to dilated cardiomyopathy if left untreated.

Multifocal atrial tachycardia occurs when multiple sites in the atria are discharging and is due to increased automaticity. It is characterised by P waves of varying morphologies and PR intervals of different lengths on the electrocardiographic trace. The ventricular rate is irregular. It can be distinguished from atrial fibrillation by an isoelectric baseline between the P waves. It is typically seen in association with chronic pulmonary disease. Other causes include hypoxia or digoxin toxicity.

Atrial tachycardia with atrioventricular block is typically seen with digoxin toxicity. The ventricular rhythm is usually regular but may be irregular if atrioventricular block is variable. Although often referred to as "paroxysmal atrial tachycardia with block" this arrhythmia is usually sustained.

Multifocal atrial fibrillation

Conditions associated with atrial tachycardia

- Cardiomyopathy
- Chronic obstructive pulmonary disease
- Ischaemic heart disease
- Rheumatic heart disease
- Sick sinus syndrome
- Digoxin toxicity

Atrial tachycardia with 2:1 block in patient with digoxin toxicity

The ABC of clinical electrocardiography is edited by Francis Morris, consultant in emergency medicine at the Northern General Hospital, Sheffield; June Edhouse, consultant in emergency medicine, Stepping Hill Hospital, Stockport; William J Brady, associate professor, programme director, and vice chair, department of emergency medicine, University of Virginia, Charlottesville, VA, USA; and John Camm, professor of clinical cardiology, St George's Hospital Medical School, London. The series will be published as a book in the summer.

Steve Goodacre is health services research fellow in the accident and emergency department at the Northern General Hospital, Sheffield; Richard Irons is consultant in accident and emergency medicine at the Princess of Wales Hospital, Bridgend.

BMJ 2002;324:594-7

A paper that changed my practice

Communication in the doctor-patient relationship

In 1972 J P Recordon, a Cambridgeshire general practitioner, wrote an article on communication in general practice. The article arose out of a Balint group that he had been attending for three years under the leadership of Dr Marie Singer.

Balint groups, organised along the lines suggested by Michael Balint, were all the rage in the early 1970s; some tended a little too much towards amateur psychoanalysis, but they certainly pioneered the examination of the nature of the consultation in general practice and analysed what subconscious undercurrents might be going on.

It was Recordon's view that it is quite all right to become emotionally involved with patients, and, indeed, that this should be encouraged—as long as enough detachment is maintained to ensure that the doctor is in control of the situation. This view confirmed what I had intuitively felt but was nervous to express, but is able to use these emotions to the greater benefit of the patient. A lot of his article is commonsense, but written in a disarming and captivating manner.

Three years later, I can confirm all his findings, and I am grateful to him for endorsing my own feelings on allowing myself to "get involved" with patients and for giving me the courage to follow his example.

Selwyn Goodacre, general practitioner, Sandlinestone, Derbyshire

What is already known on this topic

Endoscopy is a commonly used investigation for upper gastrointestinal symptoms, but its effectiveness has been questioned.

Non-invasive testing for Helicobacter pylori has been shown to predict endoscopic diagnosis in patients with dyspepsia.

What this study adds

In patients less than 55 years of age with uncomplicated dyspepsia, non-invasive testing for H. pylori is as effective and as safe as endoscopy.

Non-invasive H. pylori testing is as reassuring to the patient as endoscopy and is less uncomfortable and distressing.

Conclusion

Our current study and the previous studies, therefore, all indicate that non-invasive testing for H. pylori is as effective as endoscopy in managing patients with uncomplicated upper gastrointestinal symptoms. The non-endoscopic strategy has two potential benefits. The first is that patients find the procedure of non-invasive H. pylori testing less uncomfortable and distressing than endoscopic examination. The second is that non-invasive H. pylori testing is substantially cheaper than endoscopy. For these reasons, non-invasive H. pylori testing seems to be the preferred investigation for patients with uncomplicated dyspepsia.

Finally, it should be emphasised that our study provides information on the relative merits of only two investigational strategies. It is likely that other approaches, such as empirical treatment without investigation or the use of other investigations, will be more appropriate for certain patients.

Contributors: see bmj.com

Funding: The study was commissioned and funded by the NHS Executive research and development technology assessment programme. The views expressed in this paper are those of the authors and not necessarily those of the Department of Health.

Competing interests: None declared.

8 Garratt AM, Ruta DA, Abdalla MI, Buckingham JK, Russell IT. The SF36 health survey questionnaire: an outcome measure suitable for routine use within the NHS. BMJ 1995;306:1440-4.

(Accepted 2 February 2002)

Corrections and clarifications

ABC of clinical electrocardiography: Atrial arrhythmias

At least one reader noticed that in this article by Steve Goodacre and Richard Irons (9 March, pp 594-7), the electrocardiogram at the top of p 597 should have been captioned “Multifocal atrial tachycardia” [not fibrillation].

Protein conjugate pneumococcal vaccines

In this editorial by Vana Spoulou and colleagues (30 March, pp 729-31), our zeal to avoid too much repetition of “protein conjugate pneumococcal vaccine” unfortunately may have led to some confusion. The first sentence of the second paragraph should have read: “The need for a strict, objective assessment of the vaccine is further enhanced by serious concerns raised recently when pneumococcal polysaccharide vaccine [not “this vaccine”] was unexpectedly found to increase the rates of pneumonia in HIV infected individuals.”

Minerva

Amar Alvity and Roger Holden, the authors of the picture story about a patient with a visual field defect that was thought to be caused by excessive upper eyelid skin but was in fact secondary to an intraocular tumour (25 February, p 494), were unhappy with the tone of the text that accompanied their picture. The problem arose because the BMJ did not receive the correct wording. The authors apologise for any offence caused and have since supplied us with the version they intended us to use; this can be read at www.bmj.com/cgi/eletters/324/7335/494#263951.