



CONGRESSO

ROMA, 25 - 27 Giugno 2026

18.00 - 19.30

CARDIOPATIE VALVOLARI ED ENDOCARDITI: UPDATE 2026

Moderatori: C. Cianfrocca, Roma - A. Posteraro, Tivoli - RM

Profilassi e terapia antibiotica, gestione dello shock settico e complicanze. Update 2026

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COI

**Negli ultimi due anni rapporti di finanziamento con i
seguenti soggetti portatori di interessi commerciali in
ambito sanitario:**

Takeda, Valneva, Q-Linea, Tillots

**Table 3** New recommendations**Recommendation****Class****Level**

Section 3. Recommendation Table 1 — Recommendations for antibiotic prophylaxis in patients with cardiovascular diseases undergoing oro-dental procedures at increased risk of infective endocarditis

General prevention measures are recommended in individuals at high and intermediate risk of IE.

I**C**

Antibiotic prophylaxis is recommended in patients with ventricular assist devices.

I**C**

Antibiotic prophylaxis may be considered in recipients of heart transplant.

IIb**C**

Populations at risk of infective endocarditis

The groups of individuals at **high risk of IE in whom antibiotic prophylaxis is recommended or should be considered include the following:**

- (i) Patients with previous IE**
- (ii) Patients with surgically implanted prosthetic valves, with transcatheter implanted prosthetic valves, and with any material used for cardiac valve repair**
- (iii) Patients with congenital heart disease (CHD) (not including isolated congenital valve abnormalities)**
- (iv) Patients with ventricular assist devices as destination therapy**

Patients at **intermediate risk of IE include those with:**

- (i) rheumatic heart disease (RHD);**
- (ii) non-rheumatic degenerative valve disease;**
- (iii) congenital valve abnormalities including bicuspid aortic valve disease;**
- (iv) cardiovascular implanted electronic devices (CIEDs);**
- (v) hypertrophic cardiomyopathy.**



Section 3. Recommendation Table 2 — Recommendations for infective endocarditis prevention in high-risk patients

Systemic antibiotic prophylaxis may be considered for high-risk patients undergoing an invasive diagnostic or therapeutic procedure of the respiratory, gastrointestinal, genitourinary tract, skin, or musculoskeletal systems.

IIb**C**



Section 3. Recommendation Table 3 — Recommendations for infective endocarditis prevention in cardiac procedures

Optimal pre-procedural aseptic measures of the site of implantation is recommended to prevent CIED

I**B**

Surgical standard aseptic measures are recommended during the insertion and manipulation of catheters in the catheterization laboratory environment.

I**C**

Antibiotic prophylaxis covering for common skin flora including *Enterococcus* spp. and *S. aureus* should be considered before TAVI and other transcatheter valvular procedures.

IIa**C**

Table 6 Prophylactic antibiotic regime for high-risk dental procedures

Situation	Antibiotic	Single-dose 30–60 min before procedure	
		Adults	Children
No allergy to penicillin or ampicillin	Amoxicillin	2 g orally	50 mg/kg orally
	Ampicillin	2 g i.m. or i.v.	50 mg/kg i.v. or i.m.
	Cefazolin or ceftriaxone	1 g i.m. or i.v.	50 mg/kg i.v. or i.m.
Allergy to penicillin or ampicillin	Cephalexin ^{a,b}	2 g orally	50 mg/kg orally
	Azithromycin or clarithromycin	500 mg orally	15 mg/kg orally
	Doxycycline	100 mg orally	<45 kg, 2.2 mg/kg orally >45 kg, 100 mg orally
	Cefazolin or ceftriaxone ^b	1 g i.m. or i.v.	50 mg/kg i.v. or i.m.

- ✓ At-risk dental procedures include **dental extractions**, **oral surgery procedures** (including periodontal surgery, implant surgery, and oral biopsies), and **dental procedures** involving manipulation of the gingival or periapical region of the teeth (including scaling and root canal procedures).
- ✓ **Implant placement procedures**, and **invasive** dental procedures on established implants, however, should be covered by **antibiotic prophylaxis** in those at high risk of IE.
- ✓ The **main target** for antibiotic prophylaxis is oral **streptococci**.

Treatment of Infective Endocarditis

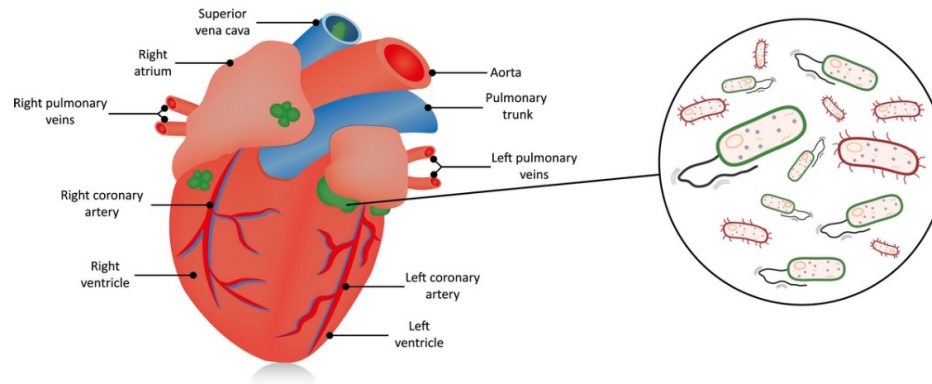
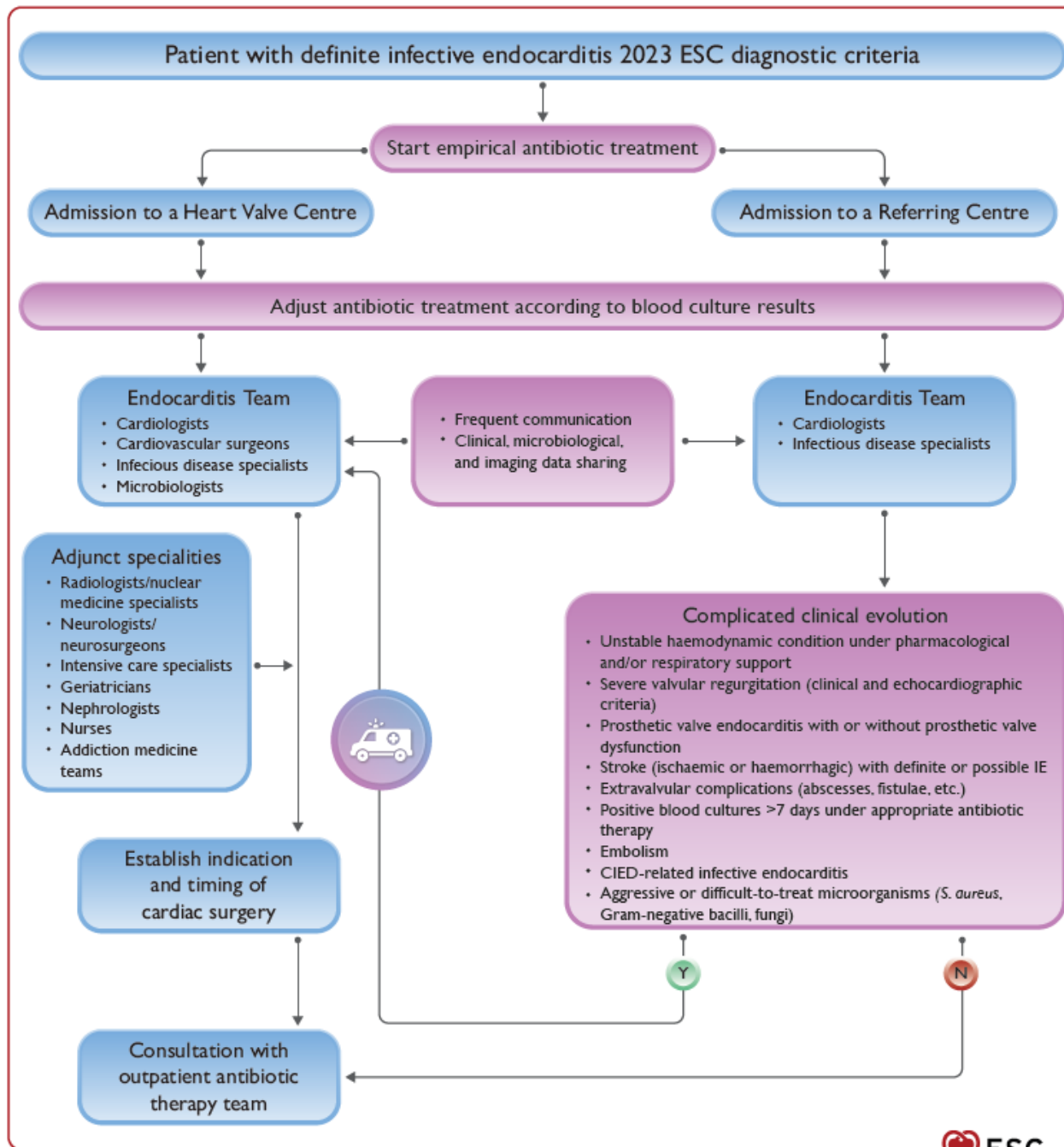


Table 7 Members of the Endocarditis Team

	Heart Valve Centre
Core members	• Cardiologists. • Cardiac imaging experts. • Cardiovascular surgeons. • Infectious disease specialist (or internal medicine specialist with expertise in infectious diseases). • Microbiologist. • Specialist in outpatient parenteral antibiotic treatment.
Adjunct specialities	• Radiologist and nuclear medicine specialist. • Pharmacologist. • Neurologist and neurosurgeon. • Nephrologist. • Anaesthesiologists. • Critical care. • Multidisciplinary addiction medicine teams. • Geriatricians. • Social worker. • Nurses. • Pathologist.

Recommendation Table 4 — Recommendations for the Endocarditis Team

Recommendations	Class ^a	Level ^b
Diagnosis and management of patients with complicated IE are recommended to be performed at an early stage in a Heart Valve Centre, with immediate surgical facilities and an 'Endocarditis Team' to improve the outcomes. ^{36–41,122,123,125,126}	I	B
For patients with uncomplicated IE managed in a Referring Centre, early and regular communication between the local and the Heart Valve Centre endocarditis teams is recommended to improve the outcomes of the patients. ^{36–41,122,123,125,126}	I	B



General principles

- ✓ Successful treatment of IE relies on **microbial eradication** by antimicrobial drugs.
- ✓ Surgery contributes by **removing infected material**.
- ✓ **Bactericidal** regimens are more effective than bacteriostatic therapy.
- ✓ **Aminoglycosides synergize** with cell wall inhibitors (i.e. beta-lactams and glycopeptides) for bactericidal activity and are useful for shortening the duration of therapy (e.g. oral streptococci) and eradicating problematic organisms. However, the side effects of aminoglycosides should be taken into consideration.

General principles

- ✓ One major hindrance to drug-induced killing is **bacterial antibiotic tolerance**.
- ✓ Tolerant microbes are not resistant (i.e. they are still susceptible to growth inhibition by the drug) but **escape drug-induced killing** and may resume growth after treatment discontinuation.
- ✓ **Slow-growing** and **dormant microbes** display phenotypic tolerance towards most antimicrobials (**except rifampin** to some extent).
- ✓ They are present in **vegetations and biofilms** (complex communities of bacteria residing within an exopolysaccharide matrix that adheres to a surface, e.g. in PVE), and justify the need for prolonged therapy to fully sterilize infected heart valves.

Phases of antibiotic treatment of infective endocarditis

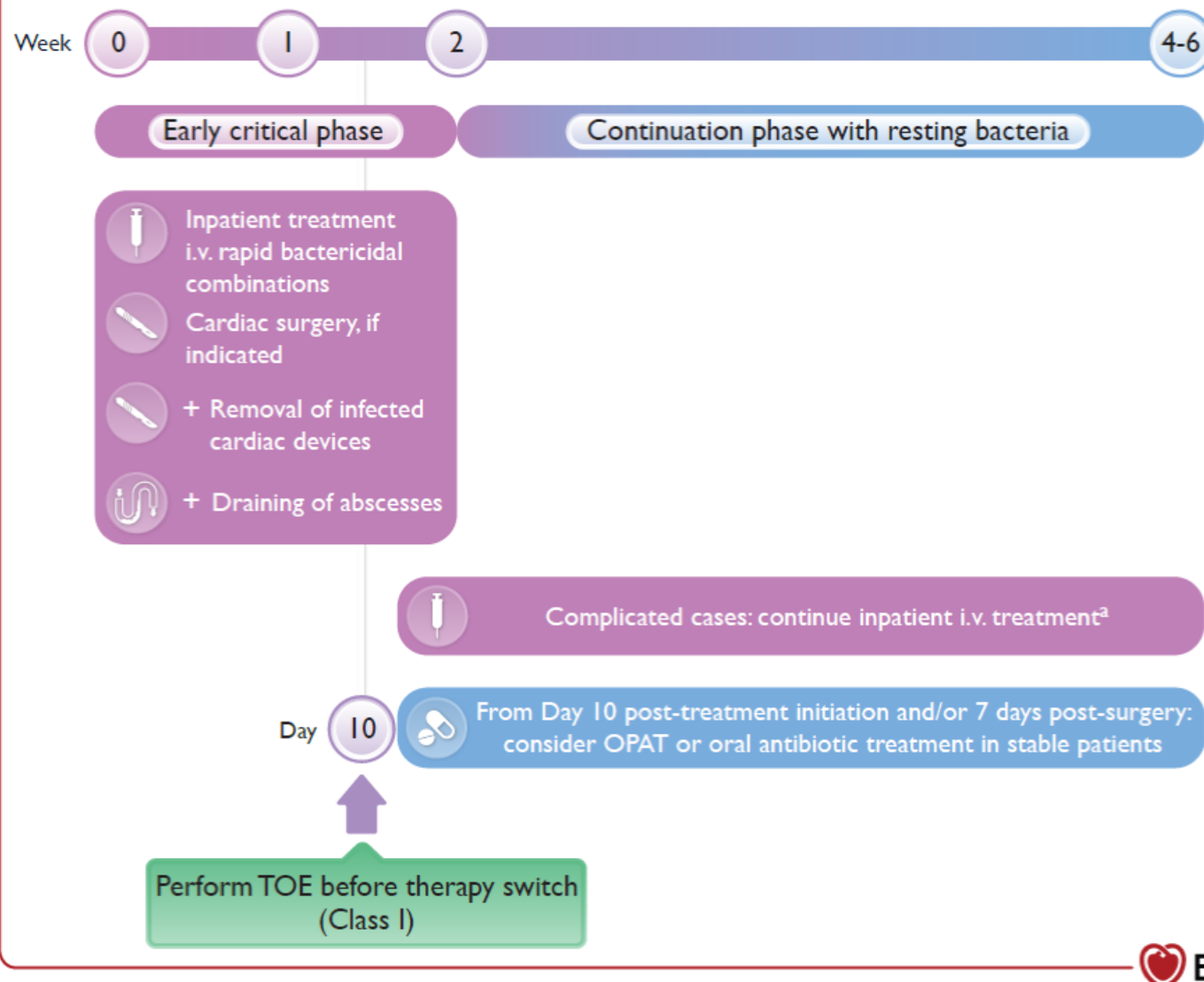


Figure 8 Phases of antibiotic treatment for infective endocarditis in relation to outpatient parenteral antibiotic therapy and partial oral endocarditis treatment. i.v., intravenous; OPAT, outpatient parenteral antibiotic treatment; TOE, transoesophageal echocardiography. ^aCriteria for switching to OPAT or partial oral treatment of endocarditis are given in the [Supplementary data online, Table S8](#).

Complications

- 1. Heart failure**
- 2. Uncontrolled infection (locally uncontrolled, septic shock)**
- 3. High risk of embolism or established embolism**



Figure 10 Proposed surgical timing for infective endocarditis. HACEK, *Haemophilus*, *Aggregatibacter*, *Cardiobacterium*, *Eikenella*, and *Kingella*; PVE, prosthetic valve endocarditis. Surgery timing: emergency, within 24 h. Urgent, within 3–5 days. Non-urgent, within same hospital admission. ^aDespite appropriate antibiotic therapy for >1 week and control of septic embolic foci. ^bE.g. patients with significant valvular dysfunction that is, or is not, a direct result of endocarditis process. ^c*S. aureus* (methicillin resistant and non-methicillin resistant), vancomycin-resistant enterococci, non-HACEK Gram-negative bacteria and fungi. ^dUrgent for *S. aureus*, non-urgent for others.

Sepsis is a syndrome defined conceptually as the development of life-threatening organ dysfunction resulting from a dysregulated host response to infection.

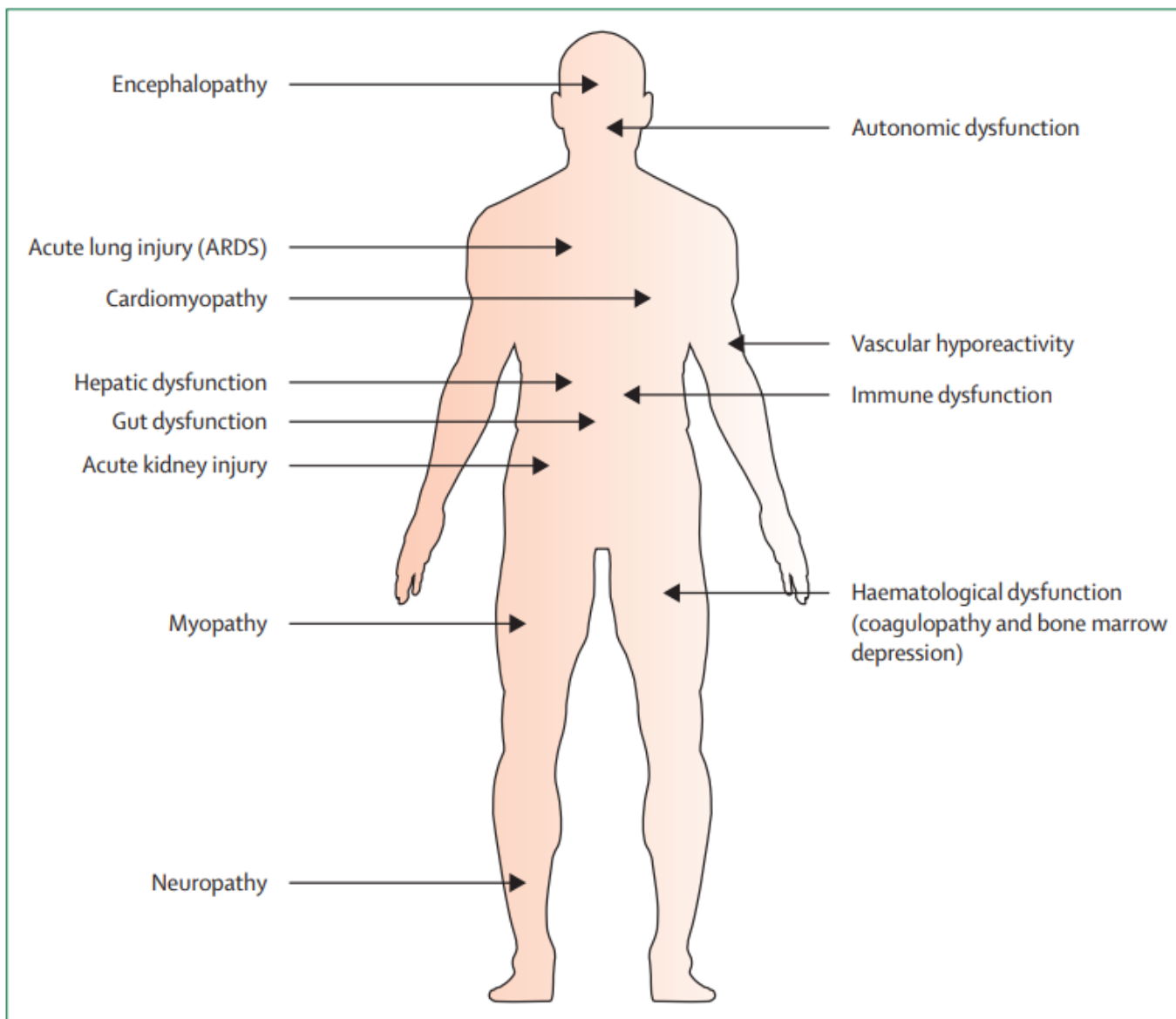


Figure 1: Organ dysfunctions in sepsis

Created in BioRender. Singer, M (2025) <https://BioRender.com/l1o1l55>. ARDS=acute respiratory distress syndrome.

Many challenges persist in clinical practice.

- **Diagnosis can be problematic, especially at first presentation and in patients with multimorbidity and/or communication issues.**
- **Infection might not be verified until positive microbiological results return.**
- **Nonetheless, causative organisms are often unidentified; therefore, infection is presumed and frequently overdiagnosed.**
- **Furthermore, acute organ dysfunction might be difficult to distinguish from pre-existing chronicity, and attributable to infection.**

The host

- Extremes of age
- Race and genetic susceptibility
- Immunosuppression
- Lifestyle
- Chronic illness
- Gut dysbiosis
- Critical illness
- Invasive devices



Pathogens

- Virulence mechanism factors
- Antimicrobial resistance patterns
- Co-infection (eg, HIV, tuberculosis, and CMV)



The healthcare system

- Few primary care providers
- Low public awareness
- Geographical barriers to access
- Resource limitations
- Workforce shortage
- Low ICU capacity
- Suboptimal quality of care
- Health care-associated infections
- Inadequate long-term follow-up



Society

- Poverty and malnutrition
- Poor sanitation and hygiene
- Inadequate vaccination coverage
- Absence of health insurance
- Underfunding of health care
- Delays in seeking health care
- Low health literacy



Other

- Low prioritisation by policy makers
- Poor implementation of national action plans
- Absence of context-specific guidelines for low-resource settings
- Challenges to doing high-quality studies
- Insufficient investment in research focused on low-resource settings

OTHER

Figure 2: Risk factors for infection and progression to sepsis

ICU=intensive care unit. CMV=cytomegalovirus.

- Sepsis is a **time-sensitive condition**: prompt identification and treatment are key.
- **No single best screening tool exists** as settings and presentations vary.
- Most approaches alert to **signs and symptoms of infection**, including markers of systemic inflammation, clinical deterioration with bedside early warning scores such as the National Early Warning Score-2 and quick Sequential Organ Failure Assessment, and illness severity (organ dysfunction and lactate).

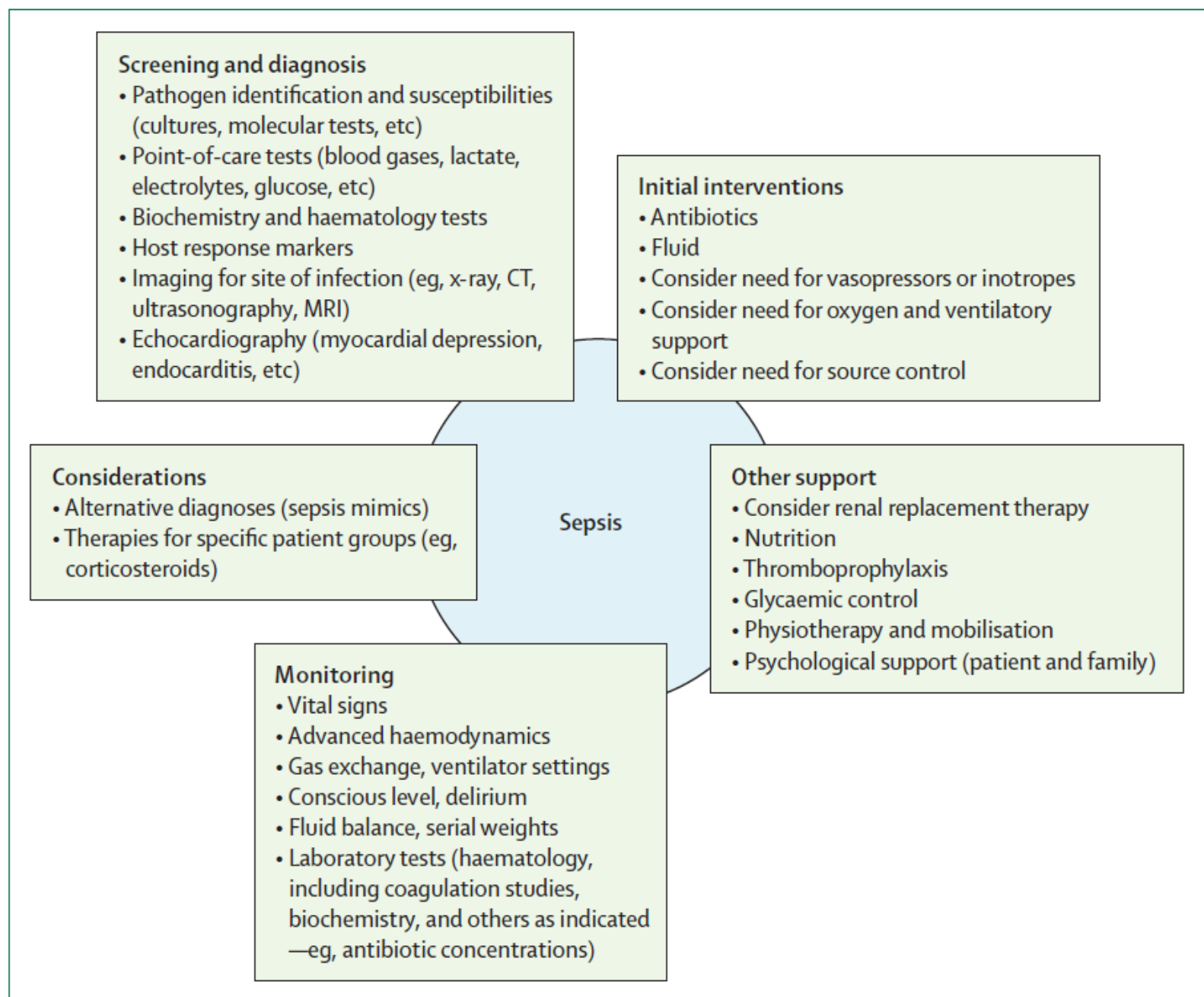


Figure 5: Sepsis management summary

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Surviving Sepsis Campaign: International Guidelines for Management of Sepsis and Septic Shock 2026

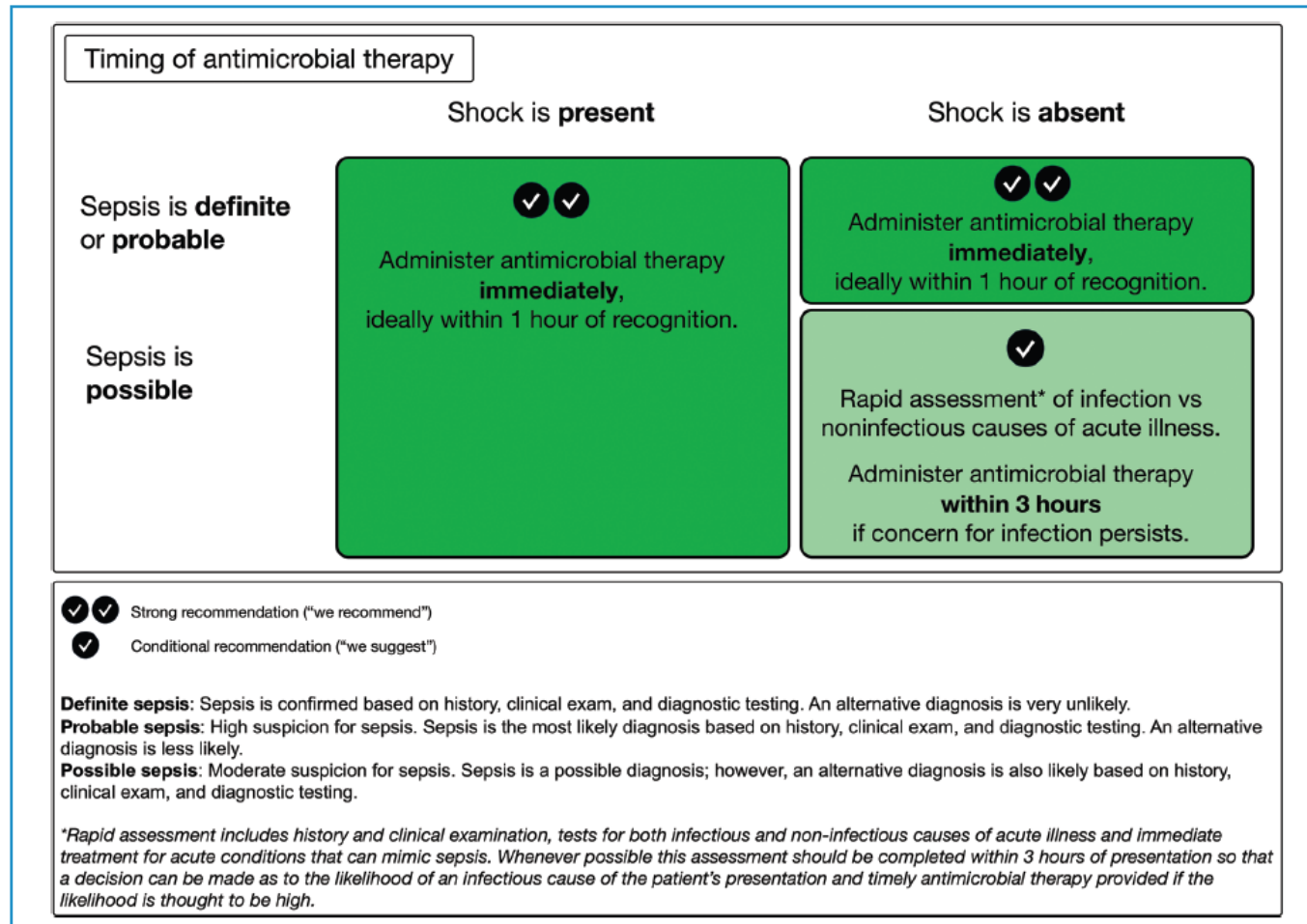


Figure 2. Antimicrobial timing framework.