

Infection-ER combine meeting

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Case Information

- * 59 y/o male
- * 1st visit day 1 12:34
- * 檢傷分級: 3級
- * 主訴: 發燒畏寒
- * GCS: E4V5M6
- * TPR: 39.9oC 心跳105/分 呼吸: 18/分
- * SpO2 96% BP 117/72 mmHg

ER course

- * 主訴: 咳嗽X 2天
- * 現病史: 咳嗽, 白痰, 左側胸口會痛, 無吐or拉肚子
- * TOCC:
 - * 旅遊史: 無
 - * 職業:建材業
 - * 無接觸史
 - * 無群聚史

ER course

- * PE:
 - * 意識: alert
 - * 頭頸: no anemic, no icteric, no inject throat
 - * 胸&心: Lt side coarse breathing sound with mild crackles, RHB, no murmur, no wheeze, no respiratory distress
 - * 腹: soft, flat, no tender or guarding, normoactive BS
 - * 四肢: freely movable, warm, no edema

ER course

- * Impression: susp pneumoniae or acute bronchitis
- * Plan: explain CXR
- * Order(12:39)
 - * CXR 2 view
 - * Tinten 1# po st
 - * Regrow 1# po st
 - * Actein 1pack po st



ER course

- * CXR findings:
 - * No pneumoniae, borderline cardiomegaly
- * Order(13:19)
 - * Recheck vital signs
 - * TPR: 37oC /101/18
 - * SpO2 95%
 - * BP 116/65 mmHg
- * 紀錄(14:07):
 - * Patient feel better, alert, coarse breathing sound, no crackle
- * Order (14:07)
 - * Tinten 1# po qid
 - * Regrow 1# po bid
 - * Actein 1pack po tid
 - * MBD & OPD follow up

ER course

- * 2nd visit
- * 檢傷分級: 2級
- * 主訴: 吃完急診藥後肢體無力
- * GCS: E4V5M6
- * 體溫: 36.4oC 心跳 91/分 呼吸: 18/分
- * SpO2 97% BP 117/85mHg

ER course

- * S: 右側無力 & numbness after discharge from ER (about 2PM), till now dysarthria noted also cough with fever, so visit ER
- * “離院後覺得怪怪的，一直冒冷汗，因一直無改善, so再度就診，目前無喘或持續咳嗽”

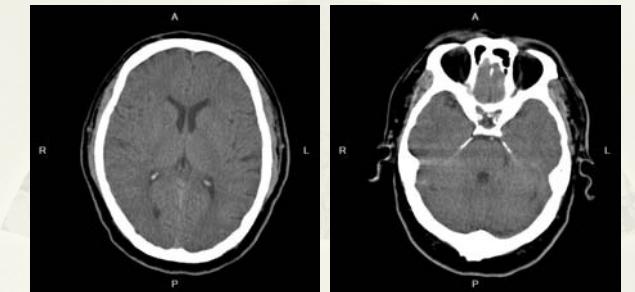
ER course

- * PE:
 - * 意識: alert, E4V5M6, slow response
 - * 頭頸: no pale conjunctiva, no icteric sclera
 - * 胸&心: clear BS, RHB
 - * 腹: soft, flat, normoactive BS, no tender or peritoneal sign
 - * 骨盆: stable
 - * 四肢: freely movable, no edema
 - * 背: unremarkable
 - * NE: no focal signs

ER course

- * Impression:
 - * 1. susp acute stroke(<3hr)
 - * 2. Acute bronchitis
- * Plan: 啟動tPA

Brain CT



ER course

- * 17: 12 Neurologist Note
 - * Impression: r/o TIA
 - * Plan:
 - * ECD/ TCD
 - * Bokey 3# po st + 1# qd
 - * Nootropil 12g iv st + qd
 - * Control BP < 220/120 and F/S <180
 - * 暫留觀 待ECD/TCD再決定MBD或admission

* 17:48轉EC20床

ER course

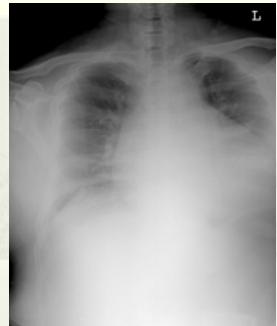
- * EC Order (18:00)
 - * Dx:
 - * 1. r/o TIA
 - * 2. acute bronchitis, r/o occult infection
 - * F/S qd /AC
 - * N/S 60ml/hr
 - * Bokey 1# po qd
 - * Nootropil 1btl iv qd
 - * If BP>220/120, or F/S >180, inform doctor

Hb	16.6	gm/dl
WBC	12.9	x1000/uL
Differential count	*****	
Segmented Neutto.	72.0	%
Lymphocyte	7.5	%
Monocyte	6.0	%
Eosinophil	0.0	%
Basophil	0.5	%
Atypical lymphocyte	0.0	%
Band	13.5	%
Metamyelocyte	0.5	%
Myelocyte	0.0	%
Promyelocyte	0.0	%
Blast	0.0	%
Nucleated RBC	0.0	/100WBC
Platelet	103	x1000/uL

Glucose	144	mg/dL
GOT(AST)	37	U/L
BUN	25	mg/dL
Creatinine	1.6	mg/dL
Na	139	meq/L
K	3.9	meq/L
eGFR	44.46	
CRP	20.700	mpg/dL
PT	14.0	second
Normal control	10.2	second
INR	1.37	Ratio
APTT	34.8	second
Normal control	32.8	second
APTT ratio	1.06	

ER course

- * 17:55
 - * U/A, B/C, Lactate
 - * 繼排ECD/TCD
- * 19:00
 - * Bedside ECHO
- * 21: 35
 - * CXR st



* 21:55

- * O2: canula 3~6L/min
- * Lasix 1amp iv st
- * U/C
- * Cefmetazole 1g iv q8h
- * Record I/O q8h

Sediment	*****
RBC	8-15 /HPF
WBC	51-100 /HPF
Epithelial cell	3-5 /HPF
Cast	Granular /LPF
cast-amount	+
Crystal	Not Found /HPF
.Cry-amount	-
Bacteria	++
Others	Mucus

Lactate 22.0 mg/dL

ER course (01/25)

- * 00:25
 - * O2 mask 6~10L/min
 - * On monitor
 - * 改IV lock
 - * Cataflam 1# po st
 - * Arrange Heart ECHO
- * 01:20
 - * Lasix 2amp iv st
- * 03:40
 - * O2 NRM 15L/min
 - * On Foley
 - * ABG (G3)
 - * Lasix 4amp iv st
 - * Millisrol 6L/min

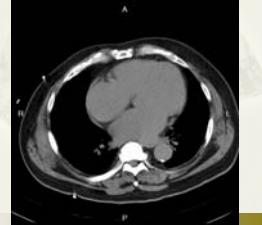
PH=7.316
PCO₂=33.9 mmHg
PO₂=367 mmHg
BE=-9 mmol/L
HCO₃=17.3 mmol/L
TCO₂=18 mmol/L
SO₂=100 %

* 03:50

- * Brain CT without contrast
- * NaHCO₃ 3amp iv st

* 04:30

- * Reconsult Neurologist
- * Admission to NRICU
- * Clexane 60mg sc st
- * Tetraspan ½ btl st
- * On critical



ER course

- * Neurologist Note(day 2 04:40)
 - * E4VAM6
 - * CN 7: R central palsy
 - * CN 8-9: dysarthria(+)
 - * Muscle power: 右: 4+; 左 5
 - * Babinski sign: 右↑/ 左↓
 - * NIHSS: 9分
- * Impression: susp Left MCA infarction, stroke evolution
- * Plan:
 - * Tetra span 0.5 btle iv st
 - * Clexane 60mg im st +q12h
 - * 轉 NR02

NR course

* Note summary:

- * HTN, CHF病史
- * 因URI症狀來ER, 回家後下午2點多 acute onset right side limbs weakness, 於是被帶回急診。
- * EKG showed Af, Septic shock & progressive右肢體無力, 4am多重做CT no ICH; 因左側MCA infarct, sepsis related住入NR

NR course

- * Impression
 - * Left MCA infarct
 - * Shock related
- * Plan
 - * Treat as septic shock
 - * Septic workup
 - * Antibiotics used
 - * Arrange Heart and Abd ECHO
 - * Supportive care

NR course

* Order(day2 05:24)

- * 輸液: N/S 40ml/hr; Tetra span 1btl iv qd
- * 抗生素: Curam 600mg iv q8h
- * Bokey 1# po qd +Clexane 60mg im q12h
- * Pantoloc 1amp q12h
- * Kerlone 0.5# po qd

WBC	10.4	×1000/μL
Differential count	-----	
Segmented Neutro.	46.0	%
Lymphocyte	6.0	%
Monocyte	6.5	%
Eosinophil	0.0	%
Basophil	0.0	%
Atypical lymphocyte	0.0	%
Band	30.0	%
Metamorphocyte	9.5	%
Myelocyte	2.0	%
Promyelocyte	0.0	%
Blast	0.0	%
Nucleated RBC	0.5	/100WBC
Platelet	48	×1000/μL
D-dimer(ELISA)	>5000	ng/mL
Lactate	118.8	mg/dL
GOT(AST)	1557	U/L
T-Bilirubin	1.9	mg/dL

* Order(09:14)

- * 抗生素:改Fortum 2g iv q8h, Clincin 600mg q8h

HBsAg	330.100	COI
Anti-HBs	2.000	IU/L
Anti-HCV	0.04	S/CO

ESR 5 mm/hr

NR course

* Heart ECHO(day 2)

- * EF: 41%
- * Rhythm : normal sinus rhythm
- * Cardiac chamber/aorta: normal wall thickness
- * Wall motion:global hypokinesis
- * Aortic valve: sclerosing change
- * Pericardial effusion: none
- * Thrombus: none
- * Others: IVC 3.07cm

NR course

* Comment:

- * Poor LV contractility, global hypokinesis
- * RA and RV dilatation, normal wall thickness
- * **Mitral valve vegetation over posterior leaflet with moderate MR**
- * Moderate TR with TRPG 20mmHg
- * IVC 3.07cm estimator RA pressure >20mmHg: **pulmonary hypertension**

* Order(day 2 16:13)

- * 抗生素加上Penicillin G 3MU q4h(Fortum, clincin)

MICU course

* Order(程Dr. service)

- * 抗生素:
 - * Aq PCN 3MU iv q6h
 - * Meropenem 1g iv q12h
 - * Teicoplanin 600Mg iv q12h X3次 then qod
- * Other Mx
 - * Dopamine run 32ml/hr
 - * Dobutamin run 25ml/hr
 - * Solu Tison (2amp in N/S 100ml run 5ml/hr)

MICU course

* Order(day 3)

- * 抗生素:
 - * Aq PCN 4MU q8h
 - * Clindamycin 900Mg q8h iv
 - * Teicoplanin 600Mg iv qod iv
- * Others
 - * Dobutamin (1mg/ml) run 25 ml/hr
 - * Levophed(4amp in 250D5W) run 10ml/hr
 - * Hydrocortisone (200Mg in N/S 100ml) run 4ml/hr

MICU course

- * OPH consultation:
- * Impression:
 - * infective endocarditis with Roth spot(OD) and SCH(OU)
 - * Cataract (OU)
 - * Conjunctival chemosis(OU)
 - * Hypertensive retinopathy Grade I (OU)
- * Abd ECHO:
 - * Fatty liver

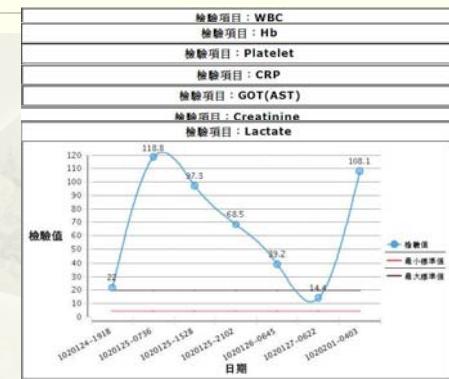
Dav 4 blood culture

PRELIMINARY BLOOD CULTURE REPORT:
 Aerobic: SALKXETJ +
 Anaerobic: SNLJ39GS +
 Two bottles of bottle set were positive cultured and final report pending.
 Microscopic finding: Gram (+) coccus in chain
FINAL BLOOD CULTURE REPORT:
 Organism:
 1 Streptococcus dysgalactiae ssp equisimilis
 Antibiotic/Culture: ST14 Streptococcus dysgalactiae ssp equisimilis
 AM CC CNZ CTX CZ E FEP LVX P VA
 S R S S R S S S S
 AM:Am(Ampicillin) CC:CC(Clindamycin) CNZ:CNZ(Cefmetazole)
 CTX:CTX(Cefotaxime) CZ:CI(Cefazolin) E:E(Erythromycin)
 FEP:FEP(Cefepime) LVX:LVX(Levofloxacin) P:P(Penicilllin)
 VA:VA(Vancomycin)

PRELIMINARY BLOOD CULTURE REPORT:
 Aerobic: SALKX7CB +
 Anaerobic: SNLJ39Q2 +
 Two bottles of bottle set were positive cultured and final report pending.
 Microscopic finding: Gram (+) coccus in chain
FINAL BLOOD CULTURE REPORT:
 Organism:
 1 Streptococcus dysgalactiae ssp equisimilis
 //
 Antibiotic/Culture: ST14 Streptococcus dysgalactiae ssp equisimilis
 AM CC CNZ CTX CZ E FEP LVX P VA
 S R S S R S S S
 AM:Am(Ampicillin) CC:CC(Clindamycin) CNZ:CNZ(Cefmetazole)
 CTX:CTX(Cefotaxime) CZ:CI(Cefazolin) E:E(Erythromycin)
 FEP:FEP(Cefepime) LVX:LVX(Levofloxacin) P:P(Penicilllin)
 VA:VA(Vancomycin)

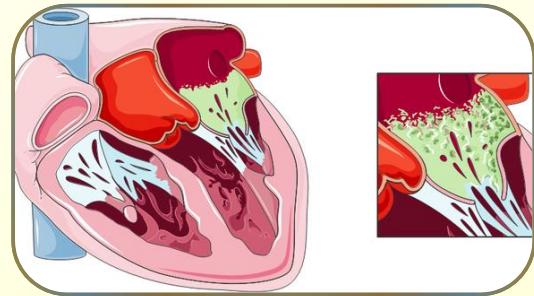
MICU course

- * CVS consultation:
 - * 開完刀可能organ failure更嚴重, 可能會出現ICH, chronic renal failure, 至少開刀前conscious要恢復, 不過開完刀conscious會變更差
- * Nephrologist consultation:
 - * Suggest H/D qd due to fluid overload



- * Day 6
 - * Try DC dormicum to monitor conscious
- * Day 7
 - * Acute cyanosis noted, ECG monitor showed Vf, s/p DC shock and CPR 100min後 家屬放棄急救

Discussion



Infective Endocarditis

Streptococcus dysgalactiae

- * Toxonomic status:
 - * *S. dysgalactiae* subsp. *equisimilis*: all b-hemolytic large colony-forming groups C and L streptococci and **human group G streptococci**
 - * 造成的疾病類似 *S. pyogenes*

Proteins with highly similar sequences	Genes or homologues described in			
	SDSE	SDSD	GAS	GBS
Fibronectin binding protein	X	X	X	
Plasminogen binding proteins	X	X	X	
Glyceraldehyde-3-phosphate dehydrogenase	X	X	X	X
Dihydrofolate reductase	X	X	X	
Protein S lytostriol binding protein	X	X	X	X
Lammin binding protein	X		X	
Streptolysin O	X	—	X	
Streptolysin S	X	—	X	
Superoxide isopeptidase (SpeA, SpeC, SpeG, SpeM, SpeN, and SpeO)	X	—	X	
Dysproteins	X			
M protein	X	IX*	X	
Capsule	X	X	X	
Cde peptide	X	—	X	X
Protein G	X	—		
Streptokinase	X	X	X	

Human Infections Due to *Streptococcus dysgalactiae*
Subspecies *equisimilis*, Clin Infect Dis. (2009) 49 (5):766-772.

Streptococcus dysgalactiae

- * 目前penicillin或其他β-lactam agents幾乎都有效
- * 因有部份susceptability下降，可考慮加上aminoglycoside
- * Aggressive infection:
 - * 加上clindamycin

Human Infections Due to *Streptococcus dysgalactiae*
Subspecies *equisimilis*, Clin Infect Dis. (2009) 49 (5):766-772.

Infective endocarditis

Table 150-2 Clinical Features of Infective Endocarditis			
Symptoms	%	Signs	%
Fever	80	Fever	90
Chills	40	Heart murmur	85
Weakness	40	New murmur	3-5
Dyspnea	40	Changing murmur	5-10
Anorexia	25	Skin manifestations	18-50
Cough	25	Osler nodes	10-23
Malaise	25	Splinter hemorrhages	15
Skin lesions	20	Petechiae	20-40
Nausea/vomiting	20	Janeway lesions	<10
Headache	20	Arterial emboli	10-50
Stroke	20	Embolic phenomena	>50
Chest pain	15	Septic complications	40
Abdominal pain	15	Mycotic aneurysm	20
Mental status change	10-15	Bimal failure	10
Back pain	10	Retinal lesions	2-10

Table 150-1 Microbiology of Infective Endocarditis (IE)

Native Valve IE (% cases)	Native valve IE (% cases)		Intracardiac Device IE (% cases)	Prosthetic valve IE (% cases)	Other devices*
	Nonaddict	IV Drug Addict			
Staphylococcal	28	68	S. aureus	23	35
Cocci/negative	9	3	Cocci/negative	17	26
Staphylococcal	21	10	Virulent group streptococci	12	8
Other streptococci	14	3	Streptococci	10	7
Enterococci species	11	4	Enterococci species	12	6
HACEK	2	0	HACEK	2	1
Fungi	1	1	Fungi	4	1
Polymicrobial	1	3	Polymicrobial	1	0
Others	4	5	Others	7	6
Culture negative	9	3	Culture negative	12	10

Abbreviation: HACEK = Haemophilus, *A*cinetobacter, *C*ardiovirulence, *E*ikenella, and *K*lebsiella group.

*Including pacemakers and implantable cardioverter defibrillators.

Clinical features from Harrison

Feature	Frequency, %
Fever	80-90
Chills and sweats	40-75
Anorexia, weight loss, malaise	25-50
Myalgias, arthralgias	15-30
Bradypnea	7-15
Heart murmur	80-85
New or worsened regurgitant murmur	20-50
Arterial emboli	20-50
Splenomegaly	15-50
Clubbing	10-20
Neurologic manifestations	20-40
Peripheral manifestations (Osler's nodes, subungual hemorrhages, Janeway lesions, Roth's spots)	2-15
Petechiae	10-40
Laboratory manifestations	
Anemia	70-90
Leukocytosis	20-30
Microscopic hematuria	30-50
Elevated erythrocyte sedimentation rate	60-90
Elevated C-reactive protein level	>90
Rheumatoid factor	50
Circulating immune complexes	65-100
Decreased serum complement	5-40

Table 150-3 Definitions of Major and Minor Criteria Used in the Duke Criteria*

Major criteria	
Positive blood culture for IE	
Typical microorganism consistent with IE from two separate blood cultures* as noted below:	
Streptococcus bovis, Viridans streptococci, HACEK group	
or	
Community-acquired Staphylococcus aureus or enterococci in the absence of a primary focus	
or	
Microorganisms consistent with IE from persistently positive blood cultures defined as:	
At least two positive cultures of blood samples drawn >12 h apart	
or	
All of three or a majority of four or more separate blood cultures (with first and last sample drawn at least 1 h apart)	
Single positive blood culture for Coxiella burnetii or antiphase I immunoglobulin G antibody titer of >1:800	
Evidence of echocardiographic involvement	
Positive ECG for IE defined as:	
Endocardial intracardiac mass on valve or supporting structures in the path of regurgitant jets, or on implanted material in the absence of an alternative anatomic explanation	
or	
Abscess	
or	
New partial dehiscence of prosthetic valve	
New valvular regurgitation (worsening or changing of preexisting murmur not sufficient)	
Minor criteria	
Predisposition: predisposing heart condition or injection drug use	
Fever: temperature >38°C (100.4°F)	
Vascular phenomena: major arterial emboli, septic pulmonary emboli, conjunctival hemorrhages, and Janeway lesions	
Immunologic phenomena: glomerulonephritis, Osler nodes, Roth spots, and rheumatoid fever	
Microbiologic evidence: positive blood culture but does not meet a major criterion as noted in Table 150-4* or serologic evidence of active infection with organism consistent with IE	
Echocardiographic minor findings were eliminated in the Modified Duke Criteria	

Streptococci

Penicillin-susceptible streptococci, <i>S. gallolyticus</i>	• Penicillin G (2–3 mL IV q4h for 4 weeks)	—
	• Ceftriaxone (2 g/d IV as a single dose for 4 weeks)	Can use ceftriaxone in patients with nonimmediate penicillin allergy
	• Vancomycin ^a (15 mg/kg IV q12h for 4 weeks)	Use vancomycin in patients with severe or immediate β -lactam allergy
	• Penicillin G (2–3 mL IV q4h) or ceftriaxone (2 g IV q6f) for 2 weeks plus	Avoid 2-week regimens when risk of aminoglycoside toxicity is increased and in prosthetic valve complicated endocarditis
	• Gentamicin ^b (3 mg/kg qd IV or IM, as a single dose ^c or divided into equal doses q8h for 2 weeks)	
Relatively penicillin-resistant streptococci	• Penicillin G (4 mL IV q6h) or ceftriaxone (2 g IV q6f) for 4 weeks plus	Penicillin alone at this dose for 6 weeks or with gentamicin during initial 2 weeks preferred for prosthetic valve endocarditis caused by streptococci with penicillin MICs of <0.1 μ g/mL.
	• Gentamicin ^b (3 mg/kg qd IV or IM, as a single dose ^c or divided into equal doses q8h for 2 weeks)	
	• Vancomycin ^c as noted above for 4 weeks	—
Moderately penicillin-resistant streptococci, nutritionally variant organisms, or <i>Gemella morbillorum</i>	• Penicillin G (4–5 mL IV q4h) or ceftriaxone (2 g IV q6f) for 6 weeks plus	Preferred for prosthetic valve endocarditis caused by streptococci with penicillin MICs of >0.1 μ g/mL.
	• Gentamicin ^b (3 mg/kg qd IV or IM as a single dose ^c or divided into equal doses q8h for 6 weeks)	
	• Vancomycin ^c as noted above for 4 weeks	—

Surgical indications

*** Absolute indication**

- * 因 valve dysfunction 引起之CHF
- * AR/MR 加上 LV/LA 壓力上升造成血行動力改變
- * Fungus 或 抗藥菌株感染
- * 出現心臟組織感染擴散的併發症

ACC/AHA Guideline Summary: Surgery for native valve endocarditis (NVE)

Class I - There is evidence and/or general agreement that surgery is indicated in patients with NVE with one of the following:
• Valve stenosis or regurgitation leading to heart failure.
• Moderate or mild regurgitation with hemodynamic evidence of elevated left ventricular end-diastolic or atrial pressures such as premature closure of the mitral valve with aortic regurgitation, rapid decelerating mitral regurgitation signal by continuous wave Doppler (v -wave cutoff sign), or moderate to severe pulmonary hypertension.
• IE due to fungal or other highly resistant organisms.
• Complications such as heart block, emboli, or aortic abscess, or destructive penetrating ulceration of the heart muscle that involves the right atrium or right ventricle, mitral leaflet perforation with IE of the aortic valve, or infection in annulus fibrosus.
Class IIa - The weight of evidence or opinion is in favor of the usefulness of surgery in patients with NVE who develop the following:
• Recurrent emboli and persistent vegetations despite appropriate antibiotic therapy.
Class IIb - The weight of evidence or opinion is less well established for the usefulness of surgery in patients with NVE who develop the following:
• Mobile vegetations larger than 10 mm with or without emboli.

Data from: Baddour LM, Casselbr EA, Chertow K, et al. ACC/AHA 2009 guidelines for the management of infective endocarditis: a joint guideline from the American College of Cardiology/American Heart Association Task Force on Practice Guidelines Writing Committee to review existing guidelines and report new guidelines for the management objectives with valvular heart disease. J Am Coll Cardiol 2009; 49(1).

Reference: Surgery of native valve endocarditis, UpToDate

https://salvagni.msu.edu.tw/content/Dentalinfo/www.uptodate.com/surgery-for-native-valve-endocarditis/source=search_result&search=surgery+endocarditis&selectedTitle=1-150

Surgery indications/timing from Harrison

Tables 124-5 Indications for Cardiac Surgical Intervention in Patients with Endocarditis

Surgery required for optimal outcome		
Moderate to severe congestive heart failure due to valve dysfunction	Emergency (same day)	Aortic or mitral regurgitation plus persistent atrial fibrillation
Partially dehisced unstable prosthetic valve		Sign of valve abscess ruptured into right heart
Persistent bacteremia despite optimal antimicrobial therapy		Rupture into pericardial sac
Lack of effective microbial therapy (e.g., fungal or Brucella endocarditis)		
<i>S. aureus</i> prosthetic valve endocarditis with an intracardiac complication	Urgent (within 1–2 days)	Value obstruction by vegetation
Relapse of prosthetic valve endocarditis after optimal antimicrobial therapy		Major embolus plus persisting large vegetation (>10 mm in diameter)
Surgery to be strongly considered for improved outcome*		
Perivalvular extension of infection		Adult aortic or mitral regurgitation with heart failure (New York Heart Association class III or IV)
Poorly responsive <i>S. aureus</i> endocarditis involving the aortic or mitral valve		Septal perforation
Large (>10-mm diameter) hypermobile vegetations with increased risk of embolism		Perivalvular extension of infection with/without new electrocardiographic conduction system changes
Persistent unexplained fever (>10 days) in culture-negative native valve endocarditis		Lack of effective antibiotic therapy
Poorly responsive or relapsed endocarditis due to highly antibiotic-resistant enterococci or gram-negative bacilli		Progressive paravalvular prosthetic regurgitation

Tables 124-6 Timing of Cardiac Surgical Intervention in Patients with Endocarditis

Timing	Indication for Surgical Intervention	Conflicting Evidence, If Any Majority of Opinions Favor Surgery
Moderate to severe congestive heart failure due to valve dysfunction	Emergency (same day)	
Partially dehisced unstable prosthetic valve		
Persistent bacteremia despite optimal antimicrobial therapy		
Lack of effective microbial therapy (e.g., fungal or Brucella endocarditis)		
<i>S. aureus</i> prosthetic valve endocarditis with an intracardiac complication	Urgent (within 1–2 days)	
Relapse of prosthetic valve endocarditis after optimal antimicrobial therapy		
Surgery to be strongly considered for improved outcome*		
Perivalvular extension of infection		
Poorly responsive <i>S. aureus</i> endocarditis involving the aortic or mitral valve		
Large (>10-mm diameter) hypermobile vegetations with increased risk of embolism		
Persistent unexplained fever (>10 days) in culture-negative native valve endocarditis		
Poorly responsive or relapsed endocarditis due to highly antibiotic-resistant enterococci or gram-negative bacilli		
Elective (unless surgically preferred)		
Progressive paravalvular prosthetic regurgitation		Staphylococcal PVE
Value dysfunction plus persisting infection after >7–10 days of antimicrobial therapy		Early PVE (<2 months after valve surgery)
Fungal (most) endocarditis		Fungal endocarditis (Candida spp.)
		Antibiotic-resistant organisms

Surgical risk in this patient

- * Population: IE + cerebral infarction
- * 目前是否需要開刀仍有爭議，因為
 - * cardiopulmonary bypass & antiocoagulants
 - 可能反而引發出血或讓腦部缺血更嚴重
- * 開刀時機:
 - * Infarction: >2 wk
 - * Hemorrhage: >4wk
 - * 因嚴重HF換valve: 考慮兩週內實行
- * 其他考量:
 - * 大vegetation(>2cm)且stroke範圍小(<2cm)要趕快開
 - 這些pt bypass 風險較小，拖久會增加recurrent emboli危險

Reference: Surgery of native valve endocarditis, UpToDate

https://salvagni.msu.edu.tw/content/Dentalinfo/www.uptodate.com/surgery-for-native-valve-endocarditis/source=search_result&search=surgery+endocarditis&selectedTitle=1-150

AHA guideline:

Prevention of infective endocarditis 2007

*** 只建議在以下患者****Table 2. Primary Reasons for Revision of the IE Prophylaxis Guidelines**

IE is much more likely to result from frequent exposure to random bacteremias associated with daily activities than from bacteremia caused by a dental, GI tract, or GU tract procedure.
Prophylaxis may prevent an exceedingly small number of cases of IE, if any, in individuals who undergo a dental, GI tract, or GU tract procedure.
The risk of antibiotic-associated adverse events exceeds the benefit, if any, from prophylactic antibiotic therapy.
Maintenance of optimal oral health and hygiene may reduce the incidence of bacteremia from daily activities and is more important than prophylactic antibiotics for a dental procedure to reduce the risk of IE.

Table 3. Cardiac Conditions Associated With the Highest Risk of Adverse Outcome From Endocarditis for Which Prophylaxis With Dental Procedures Is Reasonable

Prosthetic cardiac valve or prosthetic material used for cardiac valve repair Previous IE Congenital heart disease ^a Unprepared cyanotic CHD, including palliative shunts and conduits Completely repaired congenital heart defect with prosthetic material or device, whether placed by surgery or by catheter intervention, during the first 6 months after the procedure ^b Repaired CHD with residual defects at the site or adjacent to the site of prosthetic patch or prosthetic device (which inhibit endothelialization) Cardiac transplantation recipients who develop cardiac valvulopathy ^c
Except for the conditions listed above, antibiotic prophylaxis is no longer recommended for any other form of CHD.
^b Prophylaxis is reasonable because endothelialization of prosthetic material occurs within 6 months after the procedure.
^c Reference: Prevention of Infective Endocarditis : Guidelines From the American Heart Association, Circulation. 2007;116:1736-1754

Dental procedure Prophylaxis

Table 5. Regimens for a Dental Procedure

Situation	Agent	Regimen: Single Dose 30 to 60 min Before Procedure	
		Adults	Children
Oral	Amoxicillin	2 g	50 mg/kg
Unable to take oral medication	Ampicillin OR Cefazolin or ceftazidime	2 g IM or IV 1 g IM or IV	50 mg/kg IM or IV 50 mg/kg IM or IV
Allergic to penicillins or ampicillin—oral	Cephalexin*† OR Clindamycin OR Azithromycin or clarithromycin	2 g 600 mg 500 mg	50 mg/kg 20 mg/kg 15 mg/kg
Allergic to penicillins or ampicillin and unable to take oral medication	Cefazolin or ceftazidime*† OR Clindamycin	1 g IM or IV 600 mg IM or IV	50 mg/kg IM or IV 20 mg/kg IM or IV

IM indicates intramuscular; IV, intravenous.

*Or other first- or second-generation oral cephalosporin in equivalent adult or pediatric dosage.

†Cephalosporins should not be used in an individual with a history of anaphylaxis, angioedema, or urticaria with penicillins or ampicillin.

Reference: Prevention of Infective Endocarditis : Guidelines From the American Heart Association, *Circulation*. 2007;116:1736-1754

Thanks for your attention!