Systemic fungal infections in neonates

P C Ng

Fungal infection in newborn babies can present as mild, superficial mucocutaneous infection or as life threatening and disseminated sepsis. There are a number of ways in which fungal infection may be acquired. The infection may present at birth as congenital illness or later in life as nosocomial acquired infection.

(i) Acquired systemic fungal infections
Despite improvement in survival rates of very low birthweight (VLBW) infants, nosocomial infection remains an important contributing factor to neonatal morbidity and mortality. The emergence of fungi as important pathogens is likely to be the result of changes in the neonatal intensive care environment and their interaction with the host.

INCIDENCE
The incidence of disseminated fungal infection has not been systematically reported, due to problems in early recognition of the non-specific clinical symptoms and signs, and difficulty in confirming the diagnosis by laboratory tests. Current serological tests are not entirely reliable and microbiological cultures from sterile body fluids may only be intermittently positive, and even positive results may be misinterpreted as colonisation or contamination. A significant proportion of cases are only diagnosed at necropsy and those cases without postmortem examination would have been missed. Based on these limitations the incidence of acquired systemic fungal infections is estimated to be 2% to 5% among the VLBW infants, with species of candida being the most frequent invaders.

FACTORS PREDISPOSING TO SYSTEMIC FUNGAL INFECTION
Factors responsible for emergence of fungal infections in modern neonatal intensive care unit include impaired host defence, aggressive neonatal intensive care procedures, and prolonged use of broad spectrum antimicrobial treatment.

(b) Intensive care procedures
Invasive therapeutic and monitoring equipment has become an integral part of modern neonatal intensive care. Transcutaneous monitoring devices and adhesive tapes frequently cause superficial skin damage which facilitates the invasion of micro-organisms. Endotracheal tubes, urinary catheters, and indwelling vascular lines all bypass the skin barrier and enhance the susceptibility to systemic fungal infection by furnishing a portal of entry directly into the bloodstream. Although intravascular blood pressure monitoring devices and arterial cannulas may cause disseminated sepsis, the central venous line used for total parenteral nutrition is by far the most important apparatus responsible for fungal septicaemia. The administration of lipid emulsion may further increase the risk of fungaemia and narcotising pulmonary vasculitis due to lipophilic fungus. In addition, it has also been suggested that prolonged hospitalisation; prolonged stay in an incubator; extended period of endotracheal intubation; the use of tracheostomies; the requirement of an intraventricular shunt; and the use of lamb's wool, sticking plaster, and transparent semipermeable dressing may increase the risk of fungal colonisation and subsequent systemic infection. Equipment such as blood pressure cuffs, thermometers, oxygen saturation probes, and other diagnostic devices that are shared between infants may also promote the incidence of cross infection.
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(c) Drugs

The suppression of the normal bacterial flora by prolonged and extensive use of broad spectrum antibiotics in VLBW infants has undoubtedly contributed to the increased incidence of fungal sepsis in sick premature infants.16 Other opportunistic fungi such as aspergillus, cryptococcus, mucor, rhizopus, saccharomyces, trichosporon beigeli, etc. are uncommon and usually occur sporadically. Pathogenic fungi which include coccidioiodes, paracoccidioides, and blastomyces very rarely give rise to neonatal infections outside their endemic areas and are not a major problem in most neonatal units.

PATHOGENS

Candida and malassezia species are the two most frequent groups of opportunistic organisms causing disseminated fungal infection in premature neonates under 1500 g.6 Other opportunistic fungi such as aspergillus, cryptococcus, mucor, rhizopus, saccharomyces, trichosporon beigeli, etc. are uncommon and usually occur sporadically. Pathogenic fungi which include coccidioiodes, paracoccidioides, and blastomyces very rarely give rise to neonatal infections outside their endemic areas and are not a major problem in most neonatal units.

CANDIDA SPP

Candida albicans is the most frequent cause of fungaemia in premature VLBW infants. Other species of candida such as Candida parapsilosis, Candida tropicalis, Candida glabrata, and Candida lusitaniae have been reported to give a clinical picture indistinguishable from that caused by C albicans.

In most preterm infants, colonisation of candida occurs in utero or during delivery17 either by swallowing and aspirating the contaminated liquor, or by physical contact with the birth canal. Skin colonisation is common and often precedes the dissemination process.18 Infants with oropharyngeal and gastrointestinal tract colonisation may rarely have invasion of the bowel mucosa with resultant fungaemia.19 Candida pulmonitis secondary to aspiration is also uncommon.11 In contrast, babies colonised with candida after 2 weeks of age usually acquired the fungus nosocomially from caretakers.17

(a) Clinical presentation

The clinical features of systemic candidiasis in neonates are often non-specific and indistinguishable from bacterial sepsis. The onset is usually insidious and the mean age of infection is 33 days.1 The infant presents with temperature instability, carbohydrate intolerance, hypotension, apnoea, bradycardia, and deteriorating respiratory function which frequently requires ventilatory support. Feed intolerance, guaiac positive stools and generalised abdominal distension (without pneumatoses intestinalis) may also occur. This spectrum of symptoms and signs may be intermittent and may even be absent in some infants.1 It seems that a septic VLBW infant who deteriorates despite antibiotic treatment should always be considered to have systemic fungal infection especially when recognisable predisposing factors are present. In contrast, those cases diagnosed at necropsy tend to have had their initial presentation at a much earlier age, with fewer predisposing factors, and have followed a fulminant downhill course.

(b) Deep organ involvement

Deep organ involvement is commonly associated with systemic candidiasis. Candida spp have a high affinity for the renal tract. Haematuria, proteinuria, and pyuria are usually present with candiduria. Acute oliguria is associated with candida pyelonephritis, renal papillary necrosis, multiple parenchymal abscesses, and obstructing fungal balls.20-22 Secondary hypertension due to renal obstruction is rare.21 Candida can infiltrate the eyes via the haemagenous route causing endophthalmitis.15 These lesions are white, cotton wool ball-like and rapidly progress to involve the vitreous humour. Early detection of these lesions is important because they can cause permanent blindness. Septic arthritis9 24 and osteomyelitis23 are uncommon manifestations. Such infants usually present with a red, warm, tender, swollen, and fluid filled joint with the affected limb typically placed in an abnormal posture. The knee is the most frequent joint to be involved. It has been suggested that catheterisation of the umbilicus colonised with candida gives rise to fungaemia and subsequent synovial infection.23 Intracardiac fungal masses and endocarditis are frequently associated with indwelling right atrial lines.5 26 The fungal masses may obstruct the venous return to the heart24 26 giving rise to persistent peripheral oedema, hepatospleno-megaly, and the superior vena cava syndrome. Multiple septic emboli can occur leading to a wider disseminaton of the infection. Pulmonary candidiasis is difficult to diagnose during life. Nodular infiltrates, focal cavitation, and progressive air space consolidation are the most frequent but non-specific radiological abnormalities observed.27 Candida may also gain entry into the bloodstream by invading the bowel mucosa.19 Fungal necrotising endococolitis is usually severe and carries a poor prognosis. Central nervous system involvement is most commonly associated with intraventricular shunts.4 24 Meningitis, ventriculitis, and multiple cerebral abscesses have been described.24 38

(c) Diagnosis and investigations

The isolation of Candida spp from blood culture is the most important investigation for establishing the diagnosis of disseminated candidiasis. Blood samples may be taken from the indwelling central line, peripheral vein or artery. Although a positive blood culture is
nearly always significant, a negative culture does not exclude the diagnosis, as up to 20%-50% may not grow the organism.\textsuperscript{6} It must be emphasised that fungus grows more slowly than bacteria and antifungal treatment must therefore not be withheld in the absence of positive culture when clinical evidence is overwhelming. Examination of a Gram stained peripheral blood buffy coat preparation is sometimes useful for rapid diagnosis.\textsuperscript{29} The presence of budding yeast and hyphae in a suprapubic urine sample is diagnostic of deep seated fungal infection. Culture from other sterile sites such as cerebrospinal fluid, ascites fluid, pleura, and tissue biopsy together with culture of indwelling catheters and long lines should be obtained when clinically indicated. Candida menigitis is difficult to diagnose. Despite an increase in the cerebrospinal fluid white cell count and protein concentration, microscopy is usually negative. Moreover, these parameters may also be increased with periventricular haemorrhage without fungal infection in the central nervous system. Surveillance cultures from ears, throat, umbilicus, and rectum may also be helpful in identifying the colonisation pattern of the individual patient.

Detection of candida antigens using various enzyme linked immunosorbent assay techniques appeared to be promising with a diagnostic sensitivity of about 70% when serial specimens are evaluated.\textsuperscript{2, 30-31} However, serum levels of candida antigen can be raised in individuals who are heavily colonised or be undetectable in those with proved systemic candidiasis. The polymerase chain reaction technique, with its capability of detecting candida DNA fragments, may in future represent an important breakthrough for the early detection of candida sepsis. Measurement of candida metabolites such as D-arabinitol and biochemical tests such as the API-20c yeast test are of limited value and not routinely used in laboratories.\textsuperscript{2}

White cell and platelet counts are often abnormal in candida septicemia. Serial measurements of C reactive protein are particularly useful in monitoring the response to treatment.\textsuperscript{9, 39} A chest radiograph may show pulmonary infiltrates and progressive air space consolidation but these changes are rather non-specific and not unique for candida pneumonia.

Specific investigations are sometimes used to locate the focus of infection. Retinal examination may reveal fluffy white lesions on retina or vitreous, interlesional and leisional strands, and diffuse vitreous haze.\textsuperscript{1} Evidence of ventriculitis with foci of cavity destruction and a network of fungal strands in the ventricles may be apparent on cranial ultrasound scan.\textsuperscript{32} Cardiac and renal ultrasound scans should be routinely performed. Intracardiac fungal infection may present as thrombi in the heart chambers and vegetation on the valves,\textsuperscript{24, 26} whereas renal involvement may manifest as enlarged kidneys with highly echogenic parenchyma and fungal balls within the dilated pelvicalyceal systems.\textsuperscript{32} Percutaneous needle aspiration of the synovial cavity is indicated when neonatal fungal arthritis is suspected.\textsuperscript{23}

(d) Treatment

Three main categories of antifungal drugs are currently used for the treatment of systemic fungal infections. They are the polyclene macrolide class compounds, the azoles, and inhibitors of RNA synthesis, the fluorinated pyrimidines.

Amphotericin B belongs to the polyclene macrolide class of drugs and is the mainstay of treatment for opportunistic fungal infection. It must be administered intravenous for systemic effect as it is not absorbed enterally. The dosage could be increased in a stepwise fashion starting from 0.25 mg/kg/day up to a maximum of 1 mg/kg/day or administered as a constant dose of 0.6 mg/kg/day throughout the treatment period. The usual recommended duration of treatment is between three to six weeks. Combining amphotericin B treatment with flucytosine will permit the use of a lower dose and this combination is particularly effective for the treatment of central nervous system candida infection.\textsuperscript{4, 5} Although the side effects are better tolerated by infants, fever, renal toxicity, gastrointestinal upset, bone marrow suppression, anaphylactic reaction, and severe electrolyte disturbances including hypokalaemia and hypomagnesaemia have been well documented. Electrolytes and renal and hepatic function must be monitored closely during the treatment period.

Preparation of amphotericin B with parenteral lipid solution has been reported to reduce nephrotoxicity, improves clinical tolerance and permits a larger daily dose to be used.\textsuperscript{33}

A new formulation of amphotericin encapsulated in liposomes is commercially available. The resultant liposomal preparation is designed to maximise the delivery of amphotericin to deep seated sites of infection. The uptake by macrophages and the transport of liposomal amphotericin to the infective areas appear to play a major part in increasing efficacy and diminishing toxicity.\textsuperscript{34, 35} Two reports indicate that liposomal amphotericin is effective in the treatment of disseminated fungal infections in VLBW infants.\textsuperscript{36, 37} A starting dose of 1.5 mg/kg/day was used and this was increased progressively over a week to a maximum of 5 mg/kg/day.\textsuperscript{36} Side effects associated with conventional amphotericin such as fever, bronchospastic reactions, and renal toxicity were significantly less,\textsuperscript{36, 37} but neonatal cholestasis with raised hepatic enzymes values and conjugated hyperbilirubinaemia has been observed.\textsuperscript{37} High dose liposomal amphotericin should be tried when the fungal infection is refractory or the baby is unable to tolerate conventional treatment.

The azole group of antifungals include the imidazoles such as miconazole, ketoconazole, and the newer triazoles such as fluconazole, itraconazole, and saperconazole. They are fungistatic agents. All triazoles show promise as broad spectrum, orally active, systemic antifungal drugs with less toxicity than the
imidazoles. Fluconazole has been successfully used with minimal adverse effects for the treatment of neonatal candida septicemia, pneumonia, osteomyelitis, and central nervous system shunt infection. Despite three every two days or even daily in subsequent weeks, the half life of fluconazole varies from 55 to 89 hours in VLBW babies. Despite good evidences of its therapeutic efficacy, treatment failure in the presence of adequate minimum inhibitory concentrations has been reported. This group of drugs should, therefore, be considered as second line treatment for infants who do not respond to the standard treatment or whose condition does not permit the use of toxic antifungal agents.

Flucytosine is a synthetic compound and has the advantages of excretion via the urinary tract and good absorption when given orally. Synergy has been demonstrated with amphotericin B. Flucytosine is not recommended to be used as a single agent because resistance is a common problem and can develop during treatment. Flucytosine is given in a dose of 100 mg/kg/day and serum drug concentrations and blood counts must be monitored at least once a week because it can induce severe bone marrow suppression. Other side effects are mild and include gastrointestinal, hepatic, and renal impairment.

Acute resuscitation with plasma expander, inotropic agents, and mechanical ventilatory support are often required in the acute phase of systemic fungal illness. Removal of the infected indwelling lines and intraventricular shunt are mandatory. Surgical drainage of subcutaneous fungal abscess can be carried out at the bedside under local anaesthesia. Consultation with the orthopaedic team will help to define the situation when surgical drainage of the septic joint and convalescent immobilisation in a plaster cast is necessary. Urinary diversion, local irrigation with amphotericin B, and guidewire fragmentation of renal fungal balls have been attempted. The indications for surgical removal of intracardiac fungal masses will depend on the response to antifungal treatment, the size and mobility of the mass, the likelihood of embolisation and its haemodynamic significance on the patient. For infants under 2000 g in weight, cardio-pulmonary bypass is technically difficult and risky. The removal of intracardiac masses has been successful with the use of an inflow occlusion technique.

(c) Prognosis
Disseminated candida infections are associated with high morbidity and mortality in neonates with a significant proportion of cases being diagnosed at necropsy. Although central nervous system involvement usually has a poor prognosis and frequent neurological sequelae, intact survivors are occasionally seen. Current mortality rates range from 18% to 50% but with a trend towards prompt and aggressive antifungal treatment, outcome should improve.

(d) Prevention
Since the colonisation of the newborn with subsequent mucocutaneous candidiasis may predispose to systemic candida infection, attempts have been made to eradicate the organism from the skin and gastrointestinal tract of at risk infants. Clotrimazole vaginal tablets and cream given to pregnant women with vaginal candidiasis have significantly reduced the number of fungal contaminated newborns. Breast feeding has no impact on the incidence of oral thrush nor has chlorhexidine alcohol spray applied to the nipples of breast feeding women. The most promising prophylactic measure is perhaps the use of oral antifungal drugs, namely nystatin and clotrimazole oral gel, which have been shown to be effective in reducing the incidence of postnatally acquired gastrointestinal fungal colonisation. Nevertheless, the success of oral antifungal prophylaxis has not been translated into prevention of systemic candida infection in the newborns. As disseminated fungal infection is becoming more frequent among preterm neonates, there must be stringent control in the selection and usage of antibiotics, judicious use of indwelling lines and expedient weaning from mechanical ventilation. These must be combined with strict implementation of infection control surveillance measures in the intensive care nursery and especially careful handwashing.

MALASSEZIA SPP
Malassezia furfur is a lipophilic yeast. It is a frequent skin commensal and may colonise up to 64% to 84% of babies in a neonatal unit. Bronchopneumonia with necrotising pulmonary vasculitis and fungaemia in infants receiving parenteral lipid therapy have been reported. M furfur cannot be recovered from routine culture medium because it has an absolute nutritional requirement for medium chain fatty acids (C_{12}-C_{24}). Lipid enriched medium (such as Sabouraud’s dextrose agar with olive oil and 0-2% Tween 80) is required for its culture. As most cases resolve quickly after parenteral lipid is stopped and indwelling venous catheter is removed, only a short course of amphotericin B is required. The prognosis is usually good if the diagnosis is made early and appropriate measures promptly implemented.

(2) Congenital fungal infections
In the United States 5% to 10% of fetal deaths each year are due to intrauterine infections. Diagnosis of fetal infection is often difficult and requires elaborate laboratory tests for isolation of the organism or demonstration of its specific nucleic acids. Congenital fungal
infection is extremely rare but has been reported with candida, blastomycoses, coccidioides, and cryptococcus.

The syndrome of congenital systemic candida infection is very different from the more common acquired infection via haematogenous dissemination. Predisposing factors such as previous exposure to broad spectrum antibiotics, total parenteral nutrition, and central venous catheterisation are typically absent. The presence of a foreign body in the mother’s genital tract such as an intrauterine contraceptive device or a cervical suture contributes to colonisation of Candida spp.52–54 Although severe candidal sepsis with pneumo-

...azar, aware of prompt treatment, satisfactory response to treatment should suggest a slow, locally invasive infection with little evidence of haematogenous dissemination, even in cases with deep tissue infection.53 Congenital candidiasis typically presents in the first week of life and may simulate congenital infection of bacterial origin or other conditions such as respiratory distress syndrome or idiopathic persistent pulmonary hypertension of the newborn. In most cases, candida can be recovered from surface swabs, endotracheal or gastric aspirates. Histological examination and culture of the placenta and umbilical cord may provide important evidences for the diagnosis.2 Blood and cerebrospinal fluid specimens, however, frequently fail to demonstrate the infection despite the presence of serious generalised candidiasis.55 Antifungal treatments are similar to those used in postnatally acquired systemic candidiasis. Congenitally infected infants have a high mortality.53

Summary
Systemic fungal infections, previously considered to be a rare complication, are now frequently diagnosed in VLBW infants receiving intensive care. Confirming the diagnosis by laboratory tests is difficult and a high index of suspicion is required. Prompt and aggressive use of antifungal treatment is justified in a clinically septic neonate, especially those with a raised serum concentration of C reactive protein, who do not show a satisfactory response to antibiotics. The newer generation of liposomal amphotericin and azole antifungal drugs appear to be safe, effective, and well tolerated. With increasing awareness, prompt treatment, and better neonatal intensive care, the outcome of systemic fungal infection in preterm infants should improve.

1 Balej JE, Kliegman RM, Fanaroff AA. Disseminated fungal infections in very low birth weight infants: clinical mani-

5 Balej JE, Annable WL, Kliegman RM. Candida endoph-

14 Dunker WM, Spector SA, Fierer J, Davis CE. Malassezia fungemia in neonates and adults: complication of hyperal-

16 Bone HR, Lober RJ, Cline MJ. '3'-8-cyclic adenosine

monophosphate in the human leukocyte: synthesis, degra-

18 Fex RG, Kowarik SM, Shaw TR, Johnson RV. Mucocutaneous and invasive candidiasis among very low birth weight (μ1500 g) infants in intensive care nurs-

22 Bergman KA, Mein JP, Horrevoets AM, Montens L. Acute renal failure in a neonate due to pelvureteric candidal bezoar successfully treated with long-term systemic flu-

28 Fair RG. Systemic candida infections in intensive care nurs-

eries: high incidence of central nervous system involve-

29 Cattermole HEJ, Rivers RPA. Neonatal candida sepi-

34 Lopez-Berestein G, Kasi I, Rosenberg M. Clinical phar-

36 Lackner H, Schwinger W, Urban C, et al. Liposomal amphotericin B (AmBisome) for treatment of dissemi-

39 Cattermole HEJ, Rivers RPA. Neonatal candida sepi-


