

NUH Guide to the Clinical Management of Chicken Pox in Adults

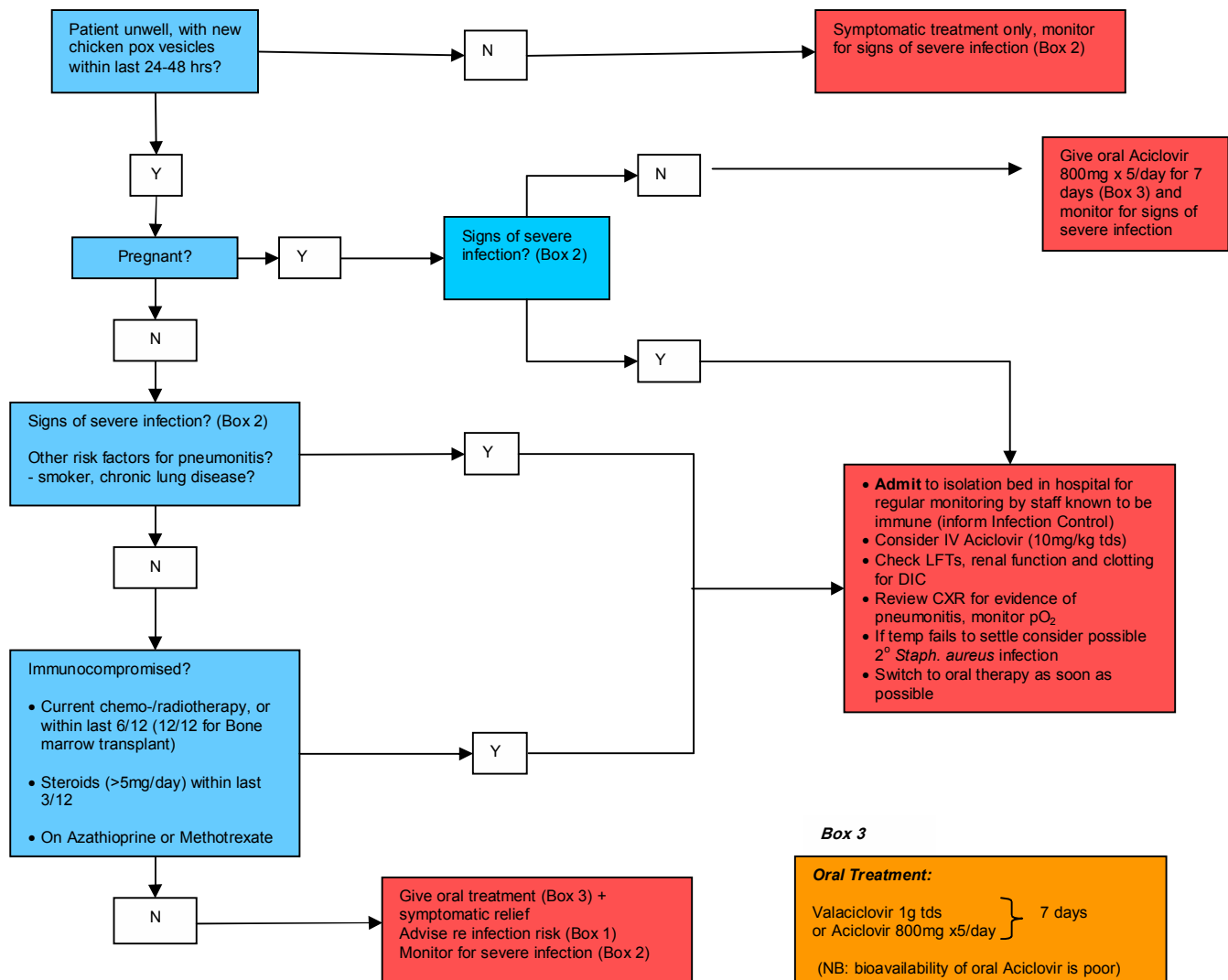
Box 1

- Chicken pox is the primary systemic infection with *Varicella-Zoster virus* (VZV)
- Acute systemic VZV has increased mortality and morbidity in adolescents and adults compared to children¹. Immunocompromised adults and non-immune pregnant women are at particular risk²
- Prompt treatment with Aciclovir reduces duration and severity of symptoms³
- There is no evidence of benefit of Aciclovir once the rash has been established for >48hrs³
- Infectivity: 2/7 prior to onset rash, until all vesicles crusted. Immunity in contacts can be assumed if clear history of clinical chicken pox
- Incubation of chicken pox: 8-21 days

Box 2

Signs of severe infection include:

- Respiratory symptoms (clinical resp signs often absent)
- Densely cropping vesicles
- Haemorrhagic rash
- Bleeding
- Any neurological changes
- Persisting fever with new vesicles >6 days after onset



Box 3

Oral Treatment:

Valaciclovir 1g tds
or Aciclovir 800mg x5/day } 7 days

(NB: bioavailability of oral Aciclovir is poor)

Intravenous treatment:

Aciclovir 10mg/kg tds

Renal impairment: dose reduction required for all forms of Aciclovir⁵

Pregnancy: No adverse data for use of Aciclovir, data inadequate for Valaciclovir⁵

References:

1. United Kingdom Advisory Group on Chickenpox. Consensus guidelines for management of varicella-zoster infection. J. Infection 1998; 36 Suppl 1: 1-83
2. Scientific Advisory Committee of Royal College of Obstetricians and Gynaecologists. Guidelines on chickenpox in pregnancy. 2001; Guideline No. 13: 1-8
3. Wallace MR, Bowler WA, Murray NB, Brodine SK, Oldfield EC3rd Treatment of adult varicella with oral aciclovir. A randomised, placebo-controlled trial. Ann Intern Med. 1992 Sep 1;117(5):358-63
4. Miller E, Marshall R, Vurdien J. Epidemiology, outcome and control of varicella-zoster infection. Reviews in Med Micro 1993; 4: 222-30
5. Joint Formulary Committee. British National Formulary, 52 ed. London: British Medical Association and Royal Pharmaceutical Society of Great Britain; Sept 06