Supplementary Material

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Research Steering Committee

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ATTIRE Site Investigators

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ATTIRE Clinical Trial Sites

Basildon, Basingstoke, Berkshire, Birmingham, Blackpool, Bournemouth, Bristol, Brighton, Coventry, Derby, Durham, Glasgow RI, Glasgow QE, Glasgow RA, Gloucestershire, Heartlands, Hull, Leeds, Liverpool, Manchester, Newcastle, North Tees, North Tyneside, Nottingham, Oxford, Plymouth, Portsmouth, Royal Free, Royal London, South Tyneside, Southampton, Surrey, Swansea, Whittington and Wigan.

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Trial Conduct

ATTIRE was conducted and reported according to the protocol, the Medicines for Human Use (Clinical Trials) Regulations 2004, (amended 2006), the European Union Clinical Trials Directive (2001/20/EC) guidelines, the principles of the International Conference on Harmonisation Good Clinical Practice under oversight of the University College London Comprehensive Clinical Trials Unit (UCL CCTU) and provisions of the Declaration of Helsinki.

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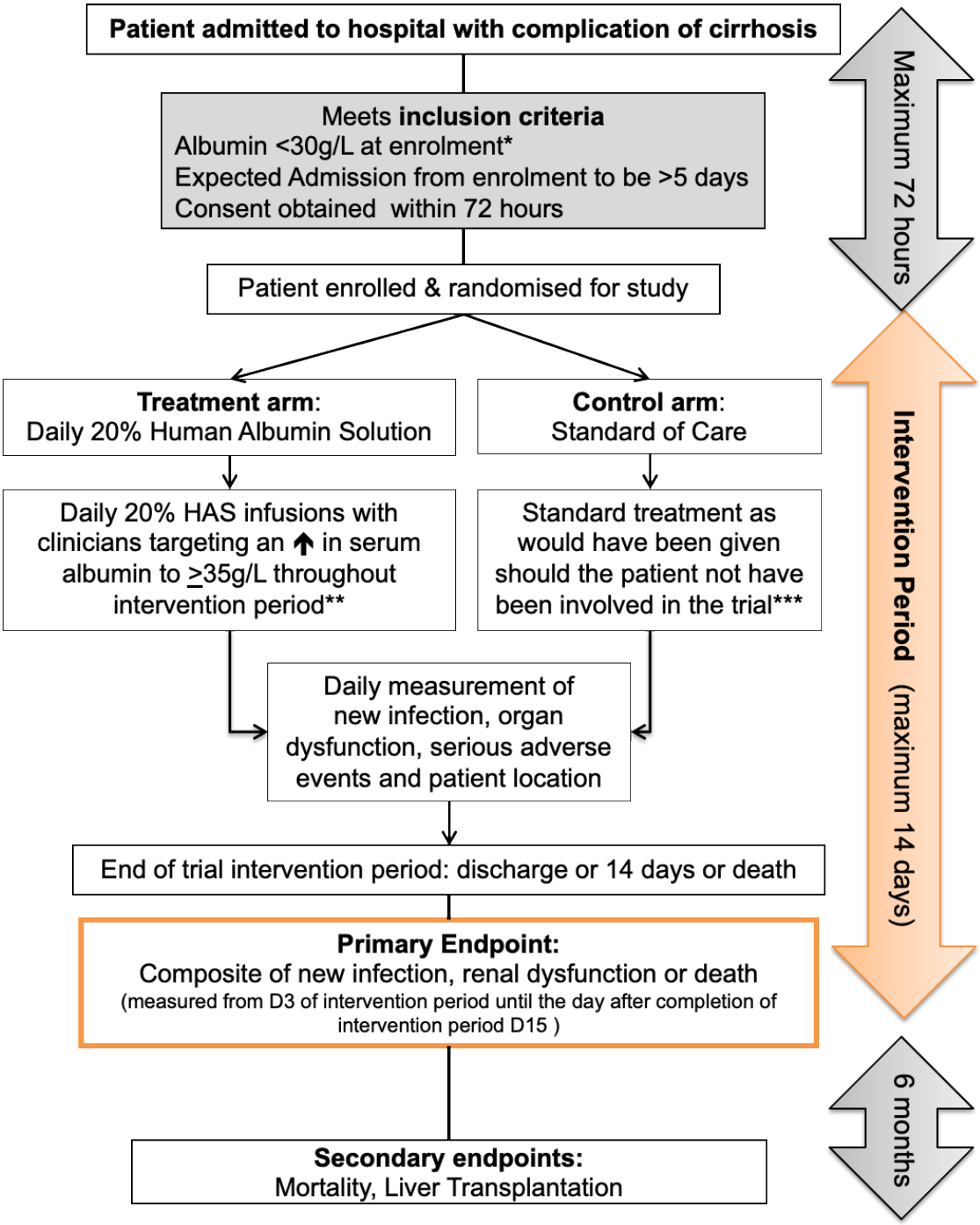
ATTIRE Study Protocol



* Which could be any time point on days 1-3 (72 hours) of admission.

** See table S2 for infusion protocol. The study aim was for treatment arm patients to achieve and sustain a serum albumin ≥30g/L. This was achieved by asking site clinicians to target a serum albumin of ≥35g/L.

*** This can only include albumin as recommended in international evidence based guidance: LVP, SBP & HRS.