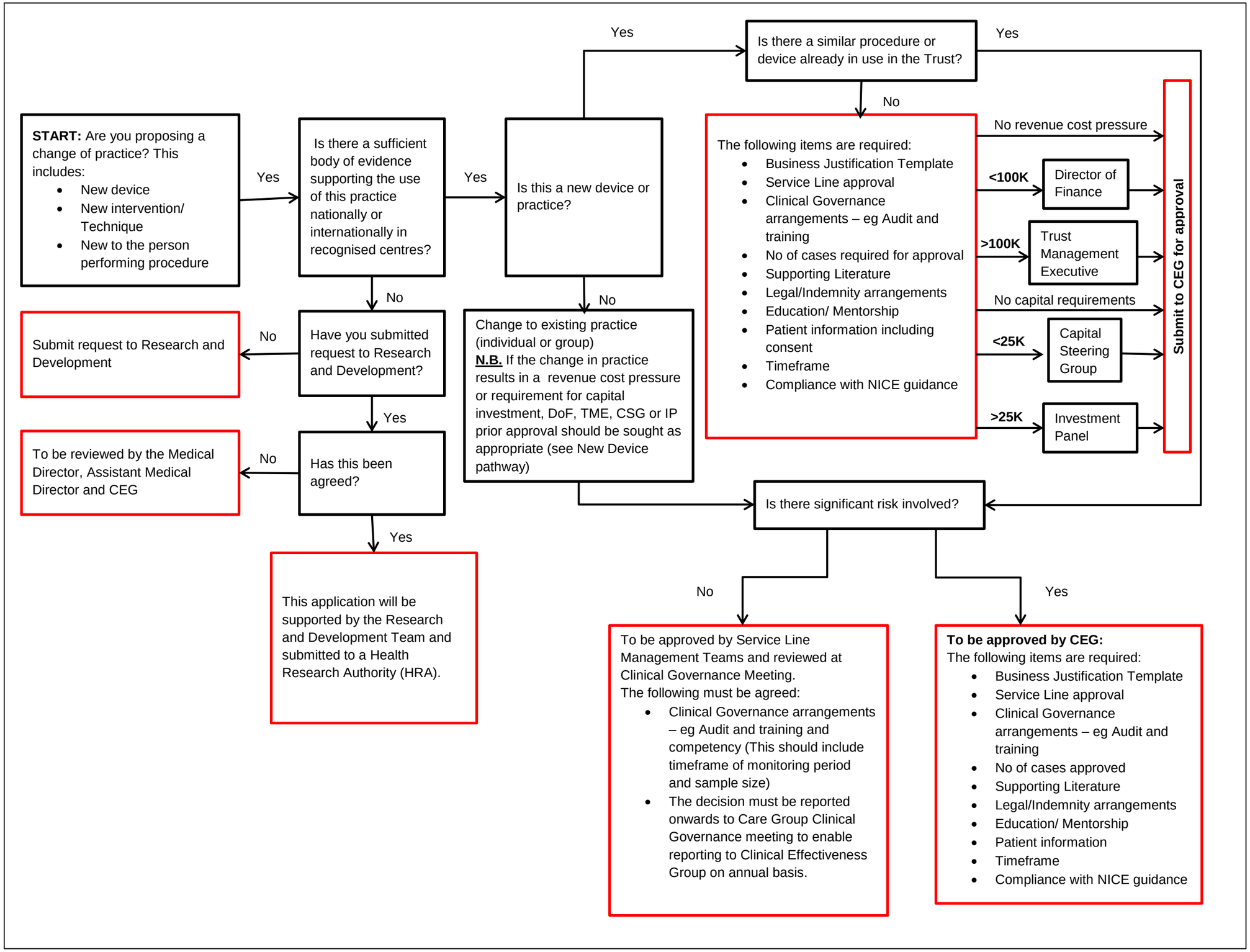
START: Are you proposing a change of practice? This includes:
New device
New intervention/ Technique
New to the person performing procedure
Yes
Is there a sufficient body of evidence supporting the use of this practice nationally or internationally in recognised centres?
Yes
Is this a new device or practice?
Yes
Is there a similar procedure or device already in use in the Trust?
Yes
No
The following items are required:
Business Justification Template
Service Line approval
Clinical Governance arrangements – eg Audit and training
No of cases required for approval
Supporting Literature
Legal/Indemnity arrangements
Education/ Mentorship
Patient information including consent
Timeframe
Compliance with NICE guidance
No revenue cost pressure
<100K
Director of Finance
>100K
Trust Management Executive
No capital requirements
<25K
Capital Steering Group
>25K
Investment Panel
Submit to CEG for approval
No
Change to existing practice (individual or group)
N.B. If the change in practice results in a revenue cost pressure or requirement for capital investment, DoF, TME, CSG or IP prior approval should be sought as appropriate (see New Device pathway)
No
Have you submitted request to Research and Development?
No
Submit request to Research and Development
Yes
Has this been agreed?
No
To be reviewed by the Medical Director, Assistant Medical Director and CEG
Yes
This application will be supported by the Research and Development Team and submitted to a Health Research Authority (HRA).
Is there significant risk involved?
No
To be approved by Service Line Management Teams and reviewed at Clinical Governance Meeting.
The following must be agreed:
Clinical Governance arrangements – eg Audit and training and competency (This should include timeframe of monitoring period and sample size)
The decision must be reported onwards to Care Group Clinical Governance meeting to enable reporting to Clinical Effectiveness Group on annual basis.
Yes
To be approved by CEG:
The following items are required:
Business Justification Template
Service Line approval
Clinical Governance arrangements – eg Audit and training
No of cases approved
Supporting Literature
Legal/Indemnity arrangements
Education/ Mentorship
Patient information
Timeframe
Compliance with NICE guidance