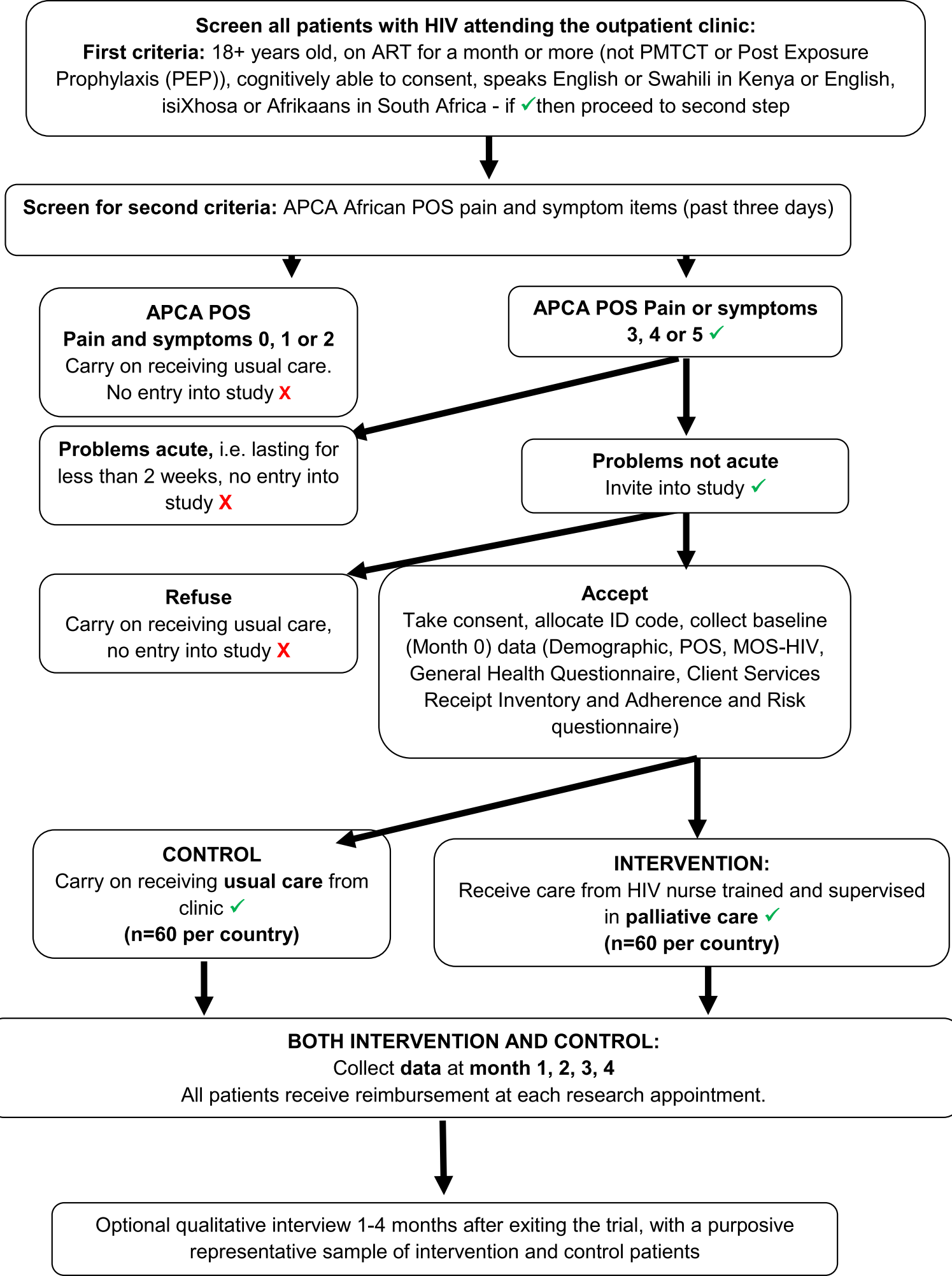
Screen all patients with HIV attending the outpatient clinic:
First criteria: 18+ years old, on ART for a month or more (not PMTCT or Post Exposure Prophylaxis (PEP)), cognitively able to consent, speaks English or Swahili in Kenya or English, isiXhosa or Afrikaans in South Africa - if ✓then proceed to second step
Screen for second criteria: APCA African POS pain and symptom items (past three days)
APCA POS
Pain and symptoms 0, 1 or 2
Carry on receiving usual care.
No entry into study X
APCA POS Pain or symptoms
3, 4 or 5 ✓
Problems acute, i.e. lasting for less than 2 weeks, no entry into study X
Problems not acute
Invite into study ✓
Refuse
Carry on receiving usual care, no entry into study X
Accept
Take consent, allocate ID code, collect baseline (Month 0) data (Demographic, POS, MOS-HIV, General Health Questionnaire, Client Services Receipt Inventory and Adherence and Risk questionnaire)
CONTROL
Carry on receiving usual care from clinic ✓
(n=60 per country)
INTERVENTION:
Receive care from HIV nurse trained and supervised in palliative care ✓
(n=60 per country)
BOTH INTERVENTION AND CONTROL:
Collect data at month 1, 2, 3, 4
All patients receive reimbursement at each research appointment.
Optional qualitative interview 1-4 months after exiting the trial, with a purposive representative sample of intervention and control patients