

Outcomes of BTK lesions and future directions

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Typical randomized device trial construct

1. The new device is compared to the existing standard of care device/surgery/medicine
2. A primary outcome endpoint is chosen not only to reflect the strengths of the new device, but also for clinical relevance
3. The endpoint will have a pre-specified time course
 - a) Occasionally the time course will be driven by number of events and therefore be unspecified
4. An expected performance level of each therapy is determined, and then a clinically relevant *delta* between them is chosen. The statistics around these assumptions will drive trial size
5. Population heterogeneity, and confounding, is minimized

Prior relevant studies



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CLI: Cutting Balloon PTA

- CTA of popliteal and infrapopliteal vessels in 73 pts with CLI
- Adjunctive stenting: 20%
- One year: no surgical bypass
- Limb salvage at 1 year: 89.5%



BTK Chill

- 115 limbs/108 patients Rutherford 4-6 treated with Cryoplasty
 - Infra-popliteal vessels between 2.5 and 5.0 mm
- Results:
 - 97% acute success
 - One-year TLR 21%
 - Overall 6 month and 1 year major amputation-free survival: 93% and 85%

| | | |
|--------|----------|------------|
| • R4: | MAmp 0% | Death: 0% |
| • R5: | MAmp 11% | Death: 0% |
| • R6: | MAmp 40% | Death: 32% |
| • +DM: | MAmp 20% | Death: 9% |
| • -DM: | MAmp 4% | Death: 11% |

BTK CHILL: Observations vis-à-vis trial design

- TLR rate acceptable, but likely restenosis rate ~40%
- Significant disparity in outcomes depending on Rutherford class, diabetes



LACI Phase 2 Registry

Laser Angioplasty for Critical Limb Ischemia

- Prospective, multi-center study
- Patients with CLI
 - Rutherford Category 4-6
- Treatment:
 - ELA of SFA, popliteal and/or infrapopliteal arteries
 - Optional adjunctive PTA and stenting
- Primary Endpoint:
 - limb salvage (freedom from amputation at or above the ankle) at 6 months



LACI 2: Descriptors

155 limbs

Rutherford Category

4

29%

5 or 6

71%

Reasons for poor surgical candidacy

Absence of venous graft

32%

Poor/no distal vessel

68%

High surgical risk

46%

Only one reason

61%

Any two reasons

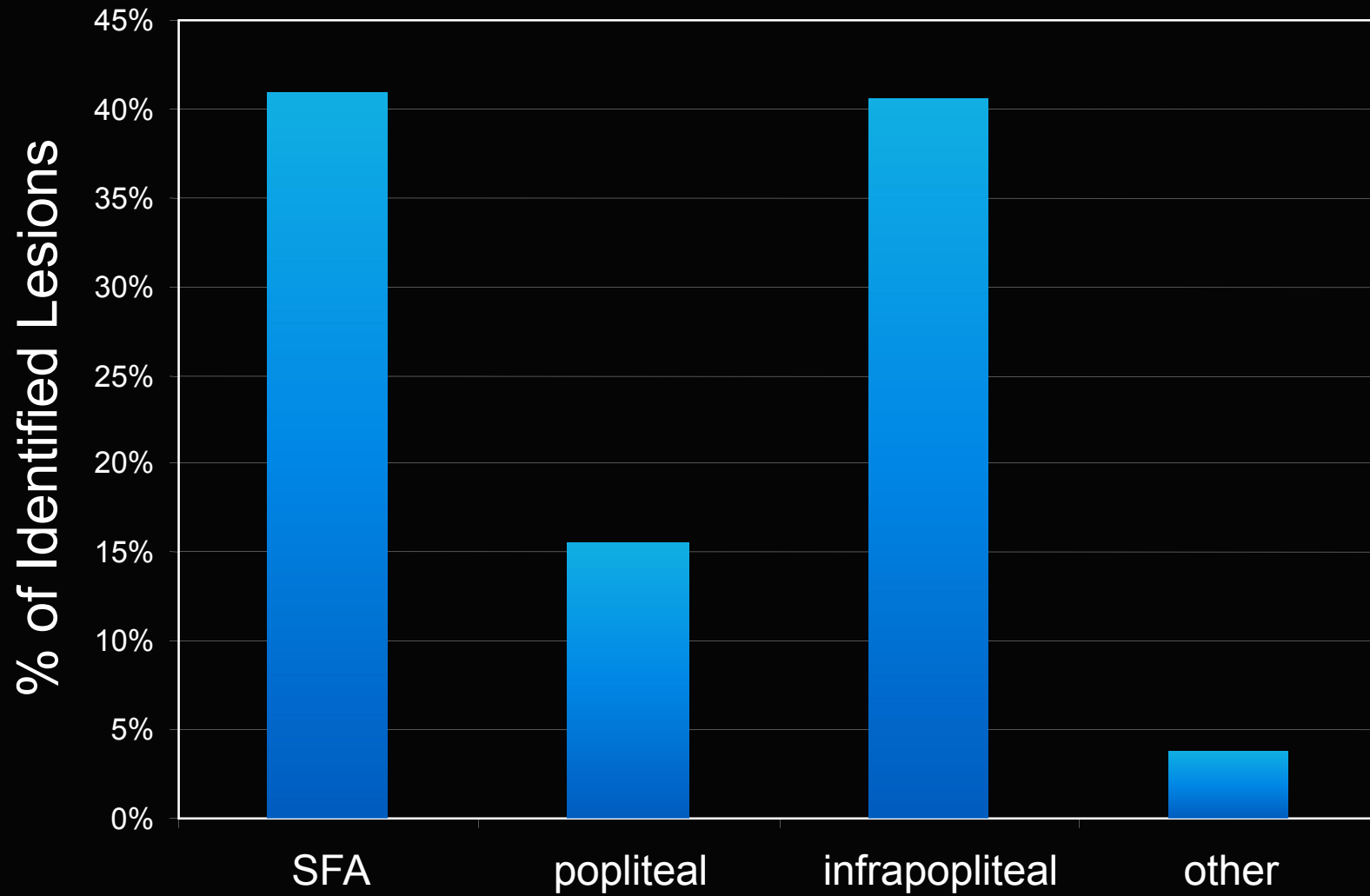
33%

All three reasons

6%



LACI 2: Vascular lesion locations



LACI 2 - Procedure Results

| | |
|--|------------|
| Guidewire crossing success | 92% |
| Laser treatment delivered | 99% |
| Adjunctive balloon | 96% |
| Stent Placement | 45% |
| Procedure Success | 85% |
| <50% residual stenosis at final | |
| Straight line flow to foot established | 89% |
| Hospital stay (days): | mean 3.0 |
| | median 1.0 |

LACI 2: 6-Month Results

| | |
|----------------------------|----------------|
| Total enrollment | 155 |
| death | 17 (11%) |
| lost to follow-up | <u>11 (7%)</u> |
| Reached 6-month follow-up | 127 |
| Major amputation | 9 (7%) |
| Survival with limb salvage | 118/127 = 93% |



LACI 2: Observations vis-à-vis trial design

- Six month outcomes non-standard time course(12 months)
- CLI represents complex disease: multiple stenoses, heterogeneous vascular distribution and occlusions
- High risk patient population with high drop-out due to mortality
- Good limb salvage rate despite this high-risk patient cohort
- Incidence of surgical intervention is very low

BASIL trial

Bypass vs. angioplasty in Severe Ischemia of the Leg

- 452 patients with CLI due to infra-popliteal disease randomized to endovascular or surgical bypass (in patients with good vein)
 - 1999-2004
 - 30 day mortality low for both
 - Surgery with more infection and MI
 - Surgery with greater 1 year costs
 - PTA TVR: 28% v. 17% at 12 months
 - No differences at 2 year but trend favoring surgery at 5 years

BASIL Results: AFS

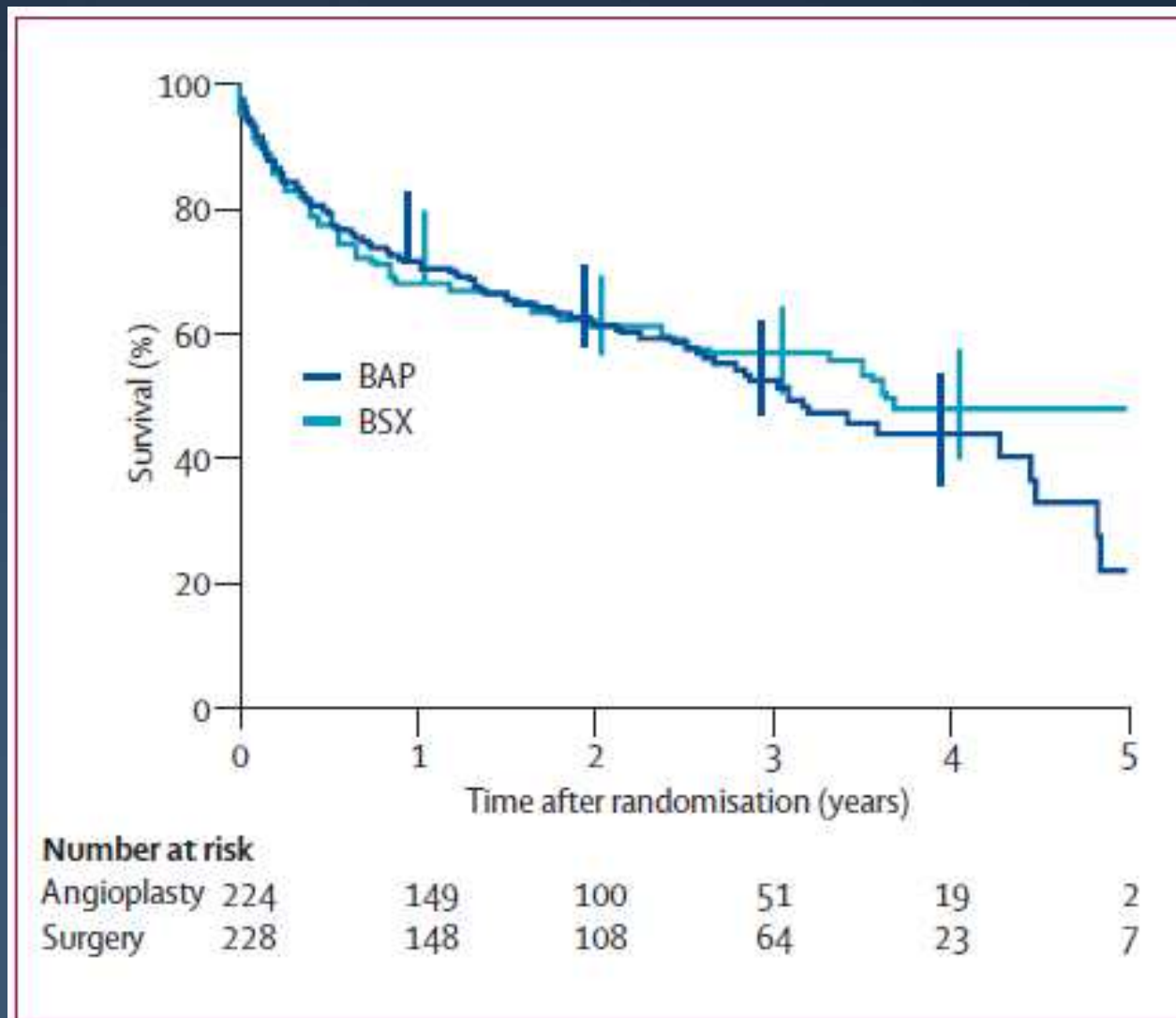


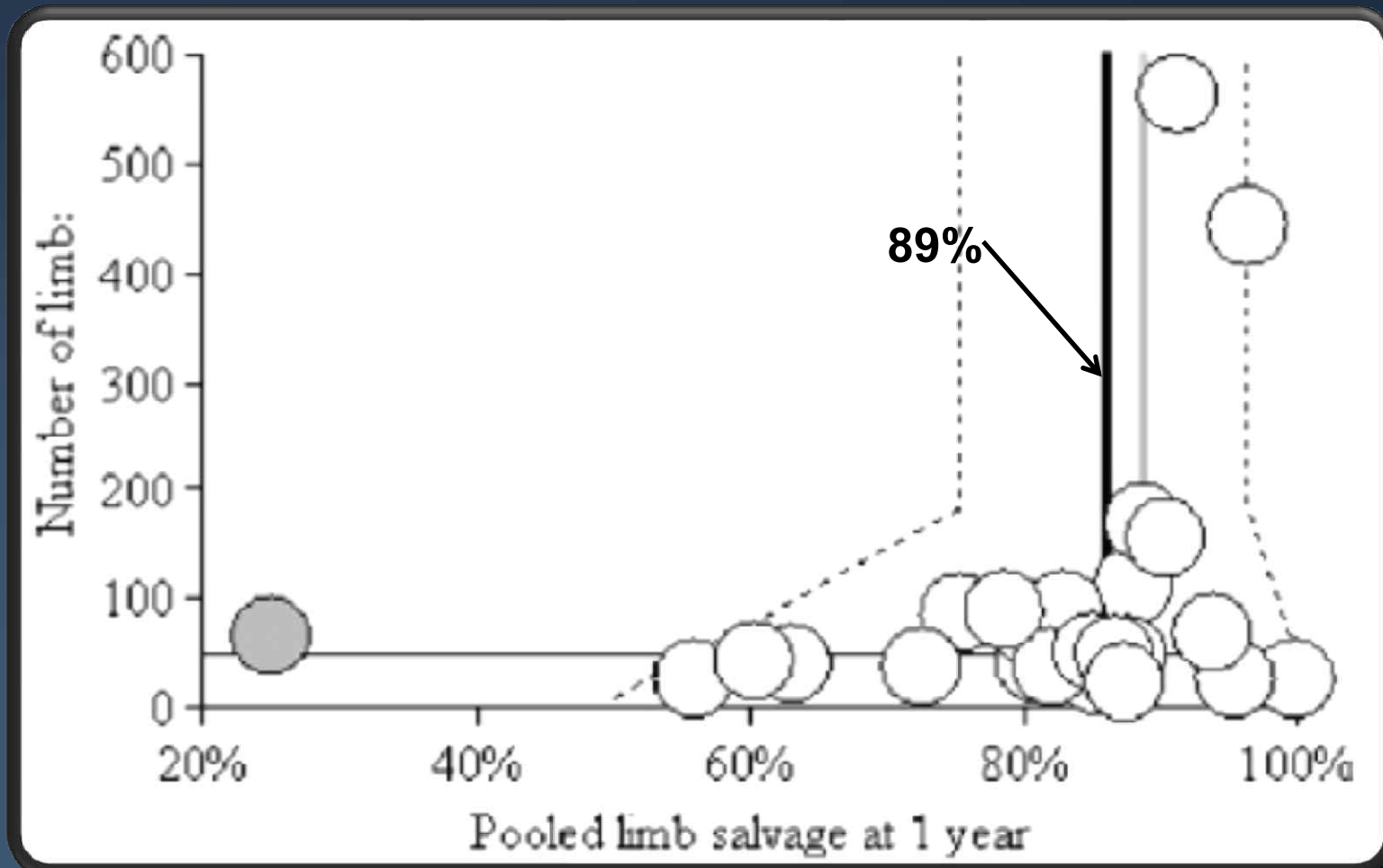
Figure 2: Amputation-free survival after bypass surgery and balloon angioplasty

Observations from BASIL

- Comparing with a surgical standard, endovascular approach to CLI is a reasonable alternative for the endpoint of limb salvage



Mets-analysis: 12 month limb-salvage



Data from meta-analysis of infra-popliteal intervention for CLI

Table II. Meta-analysis results of crural percutaneous transluminal angioplasty and popliteal-to-distal bypass^a

| Result | 1 month | 6 months | 1 year | 2 years | 3 years |
|-------------------|------------|------------|------------|------------|-------------|
| Primary patency | | | | | |
| PTA | 77.4 ± 4.1 | 65.0 ± 7.0 | 58.1 ± 4.6 | 51.3 ± 6.6 | 48.6 ± 8.0 |
| Bypass | 93.3 ± 1.1 | 85.8 ± 2.1 | 81.5 ± 2.0 | 76.8 ± 2.3 | 72.3 ± 2.7 |
| P | <.05 | <.05 | <.05 | <.05 | <.05 |
| Secondary patency | | | | | |
| PTA | 83.3 ± 1.4 | 73.8 ± 7.1 | 68.2 ± 5.9 | 63.5 ± 8.1 | 62.9 ± 11.0 |
| Bypass | 94.9 ± 1.0 | 89.3 ± 1.6 | 85.9 ± 1.9 | 81.6 ± 2.3 | 76.7 ± 2.9 |
| P | <.05 | <.05 | <.05 | | |
| Limb salvage | | | | | |
| PTA | 93.4 ± 2.3 | 88.2 ± 4.4 | 86.0 ± 2.7 | 83.8 ± 3.3 | 82.4 ± 3.4 |
| Bypass | 95.1 ± 1.2 | 90.9 ± 1.9 | 88.5 ± 2.2 | 85.2 ± 2.5 | 82.3 ± 3.0 |
| Patient survival | | | | | |
| PTA | 98.3 ± 0.7 | 92.3 ± 5.5 | 87.0 ± 2.1 | 74.3 ± 3.7 | 68.4 ± 5.5 |
| Bypass | NA | NA | NA | NA | NA |
| PTA | 98.3 ± 0.7 | 92.3 ± 5.5 | 87.0 ± 2.1 | 74.3 ± 3.7 | 68.4 ± 5.5 |
| Bypass | 98.3 ± 0.7 | 92.3 ± 5.5 | 87.0 ± 2.1 | 74.3 ± 3.7 | 68.4 ± 5.5 |

1. The new device is compared to the existing standard of care device/surgery/medicine

- The standard of care in critical limb ischemia is bypass surgery, except when it isn't:
 - Amputation is still prevalent
 - As many as 45% of patients with CLI do not have suitable ipsilateral GSV
 - The BASIL/LACI trial demonstrated both a mixed lesion location and “primitive” PTA
 - Majority of patients had SFA, 62% had infra-popliteal, PTA
 - 20% initial failure rate
 - BASIL demonstrated parity between the surgical standard, when it was available

Vascular Surgical Trends: A Changing Standard of Care

Revascularization Procedures by Vascular Surgery 2002-4

| | 2002 | 2003 | 2004 | % change |
|--------|------|------|------|----------|
| Endo | 82 | 123 | 207 | +152% |
| Bypass | 218 | 219 | 144 | -34% |



3. A primary outcome endpoint is chosen not only to reflect the strengths of the new device, but also for clinical relevance

The most relevant clinical endpoint is amputation-free survival/limb salvage, but does not highlight the strengths of a device which improves patency

3a. The endpoint has a pre-specified time course

A 1-year time course appears to be most appropriate

- Although this may not be long enough to highlight a patency advantage



4. An expected performance level of each therapy is determined, and then a clinically relevant *delta* between them is chosen. The statistics around these assumptions will drive trial size

- Problem #1: Endovascular *limb-salvage* rates are not significantly differentiated between therapies thus far
- Problem #2: Endovascular *patency* data is limited, but suggests that the relationship to limb-salvage is only moderate



5. Population heterogeneity, and confounding, is minimized

- Inclusion of Rutherford classes 4-6 leads to heterogeneity in outcomes
 - As demonstrated in LACI 2
- Both LACI and BASIL demonstrated significant lesion location heterogeneity
- Even assuming intervention is limited to infra-popliteal vessels, considerable variability in patterns of disease exist



Patterns of infra-popliteal anatomy in CLI: what to allow in studies?

- Stenosis/occlusion of the distal popliteal/TP trunk
- Stenosis of multiple vessels
- Occlusions of 1 or 2 vessels with diseased remaining vessel to foot
 - Last remaining vessel is the peroneal which incompletely collateralizes AT/PT at the ankle
- Patent single AT or PT to the foot, but incomplete plantar arch results in ischemic dermatomes

Summary of challenges

- Evolving standard of care away from surgery
- The established primary endpoint is not well defined, not well described according to patency, and not well differentiated
- Time course of follow-up may be too short to establish value of patency
 - Possible reformation of wounds is countered by subject deaths
- Marked heterogeneity in various aspects of CLI intervention
- Above combine to make statistical assumptions less well defined, thus requiring more patients, longer trials, and making success less certain

Possible solutions

- Combine limb-salvage with another meaningful endpoint (e.g., patency, wound healing)
- Be prescriptive regarding intervention to reduce heterogeneity
 - Vessel location
 - Number of vessels
 - Specify allowed anatomy
 - Limit Rutherford class inclusions
- These will increase time course of enrollment, but should allow proof of the value of patency



Overview

- Infra-popliteal anatomy and implications
- Critical limb ischemia definitions
- Importance of limb salvage
 - Consequences of amputation
- Prior interventional results
 - Laser
 - Cryoplasty
 - BASIL
- Randomized trial design challenges

Critical limb ischemia: definitions

- Rutherford classification
 - R4: Resting symptoms
 - R5: Minor tissue loss
 - R6: Major tissue loss
- Fontaine classification
 - FIII: Resting symptoms
 - FIV: tissue loss



Rutherford 5



Prognosis after amputation

- 2 year mortality rates 40%-50% following major amputation



Overview

- Define the typical trial design for new devices
- Present representative available data on infra-popliteal therapy
- Define unique regulatory challenges based on 3 characteristics of infra-popliteal disease
 - Variability in natural history among classifications
 - Anatomic variability
 - Clinically relevant endpoints

BASIL Results: Mortality

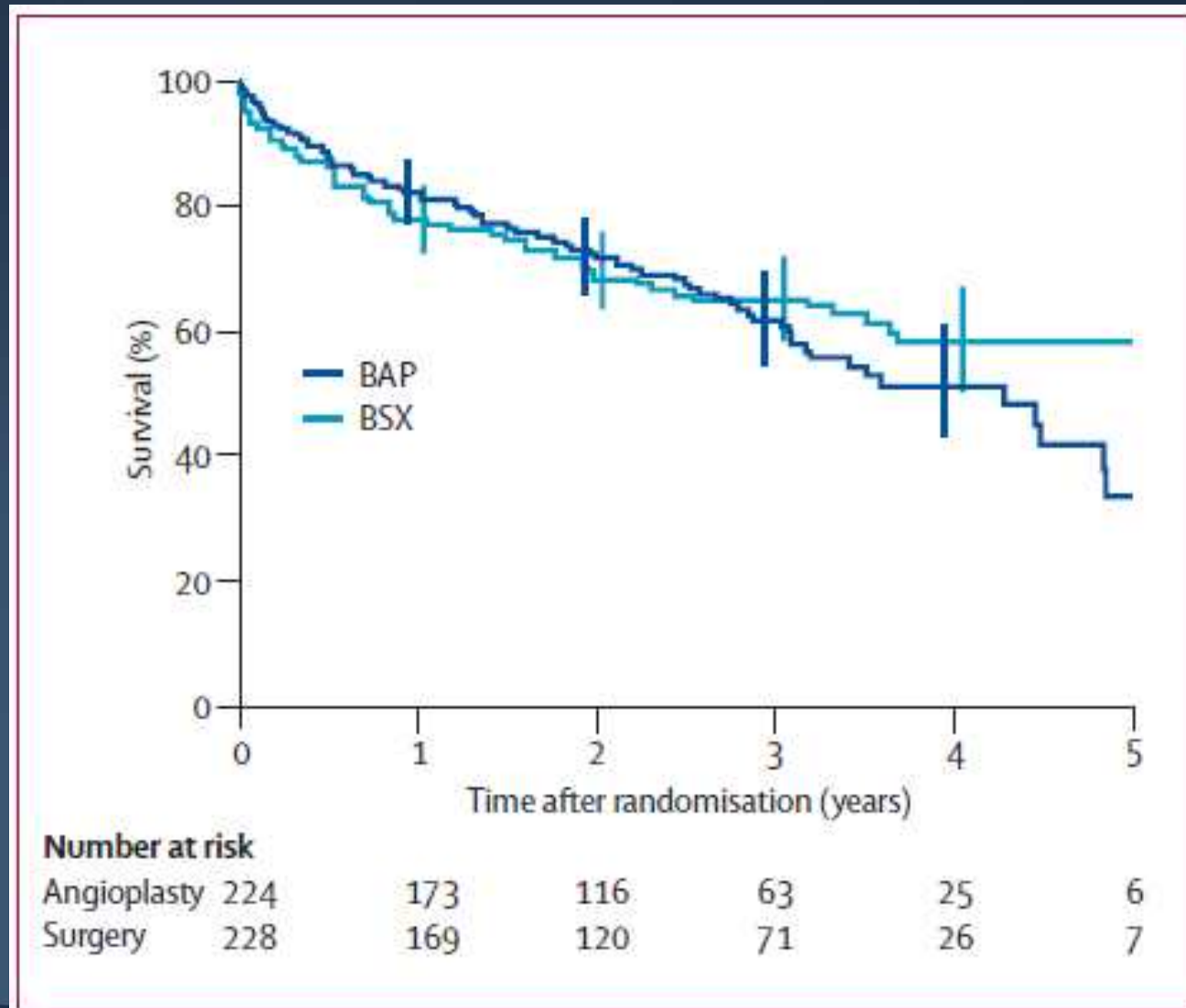


Figure 3: All-cause mortality after bypass surgery and balloon angioplasty