

Is the Palma Procedure Still Relevant in the Endovascular Era?

A Meta-Analysis of Long-Term Outcomes in Femoro-Femoral Venous Bypass

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Disclosures and Roadmap

DISCLOSURES

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Does the Palma / Palma–Dale femoro-femoral venous bypass still have a meaningful role when iliac venous stenting is the dominant therapy?

ENDOVASCULAR ERA

Iliac venous stenting is first-line for most symptomatic post-thrombotic and non-thrombotic obstruction.

- ✓ High technical success
- ✓ Low procedural morbidity
- ✓ Rapid recovery

A SUBSET REMAINS CHALLENGE

Patients in whom endovenous recanalization is impossible, anatomically unfavorable, or has already failed.

- ▶ Long-segment chronic total occlusions
- ▶ Repeat in-stent restenosis / occlusion
- ▶ Post-trauma iliac vein ligation
- ▶ Anatomically unstentable inflow

Rationale – why revisit an open operation?

44%

of post-thrombotic limbs fail first-line iliac stenting by 5 years — in-stent restenosis or re-thrombosis.

(Rollo, JVS-VL 2018; Razavi 2019)

≈15%

of chronic iliofemoral obstructive disease is anatomically un-stentable — long-segment CTO, post-trauma ligation, IVC / bilateral iliac involvement.

(Seager 2016; Texas Heart Inst J)

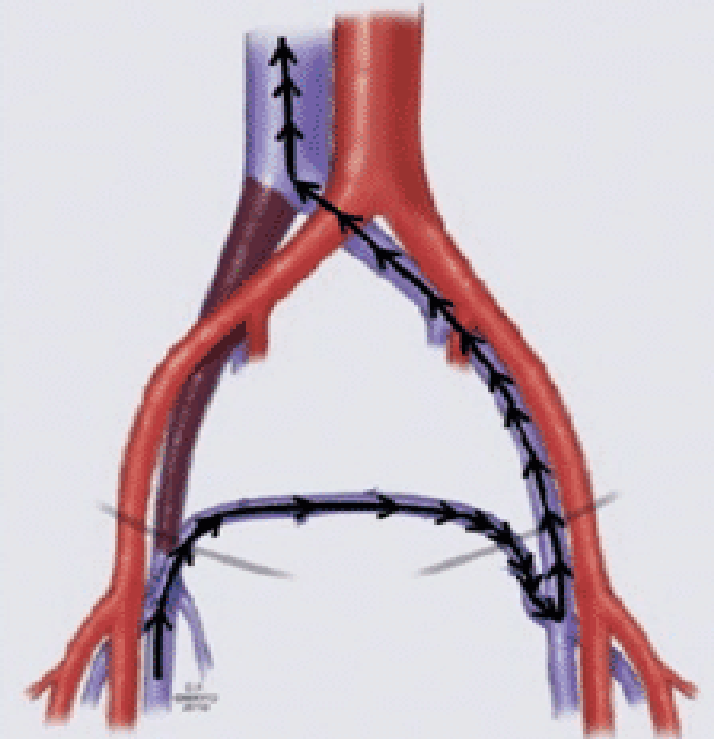
20–40%

ulcer recurrence despite successful recanalization, driving repeat endovascular re-intervention and sustained disability.

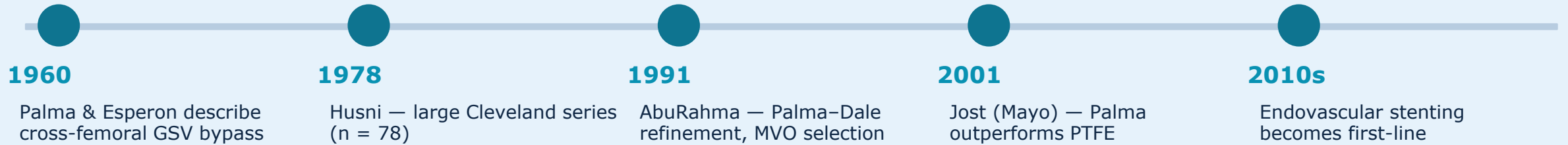
(Sarici, JVS-VL 2024)

THE UNANSWERED QUESTION

Does the Palma / Palma–Dale femoro-femoral venous bypass retain a meaningful role for the subset who cannot be served by stenting?



From Palma–Esperon (1960) to the modern stent era



THE OPERATION — in one sentence

Contralateral great saphenous vein is mobilized, tunneled suprapubically, and anastomosed to the ***common femoral vein of the obstructed limb***, restoring outflow through the patent contralateral iliac system.

Indications: chronic post-thrombotic iliofemoral obstruction · disabling May–Thurner physiology · post-trauma iliac vein ligation.

Systematic review and proportional meta-analysis

DESIGN

PRISMA 2020 systematic review of cohorts reporting Palma / Palma–Dale crossover bypass.

DATABASES

PubMed/MEDLINE · Embase · Scopus · Web of Science (inception → Dec 2025).

STATISTICS

- ▶ Logit-transformed proportional meta-analysis
- ▶ Continuity correction 0.5
- ▶ Inverse-variance fixed-effects + DerSimonian–Laird random-effects
- ▶ Heterogeneity: Cochran Q, τ^2 , I^2

OUTCOMES

Primary — graft patency at last reported follow-up. Secondary — clinical improvement, complications, mortality.

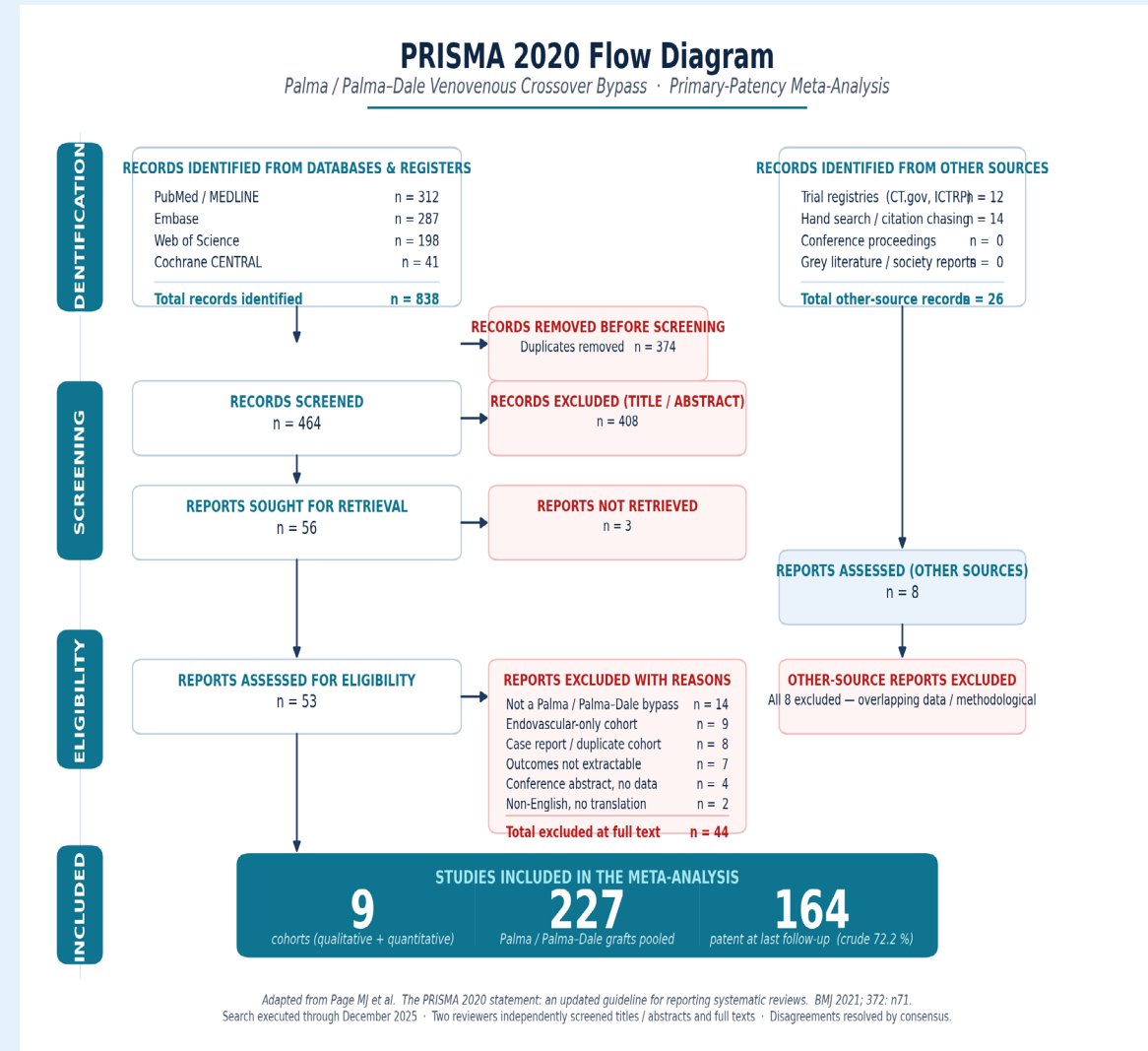


Figure . PRISMA 2020 flow.

Quality of evidence — 4 of 9 studies high quality, none at critical risk

Study	Selection /4	Comparability /2	Outcome /3	TOTAL /9	Judgement
Husni (1978)	3	1	2	6	Moderate
Hutschenreiter (1979)	2	0	2	4	Higher
AbuRahma (1991)	3	1	3	7	High
Gruss (1997)	3	1	2	6	Moderate
Jost (Mayo, 2001)	4	2	3	9	High
Alcocer (2008)	2	0	2	4	Higher
Kim (2009)	3	1	3	7	High
Garg (Mayo, 2011)	4	2	3	9	High
Kosasih (2017)	3	1	2	6	Moderate

OVERALL JUDGEMENT

9 studies

227 limbs reconstructed

4 / 9 high quality (NOS ≥ 7)

3 / 9 moderate (5–6)

2 / 9 higher concern (< 5)

No study at critical risk

Key point — modern cohorts (Jost 2001, Garg 2011, Kim 2009) had rigorous imaging-based follow-up across all three NOS domains.

Figure 2. Per-study NOS judgement across Selection, Comparability, and Outcome (adapted Newcastle–Ottawa Scale for single-arm cohorts).

INCLUDED STUDIES

9 cohorts · 227 Palma bypasses · 4 decades of evidence

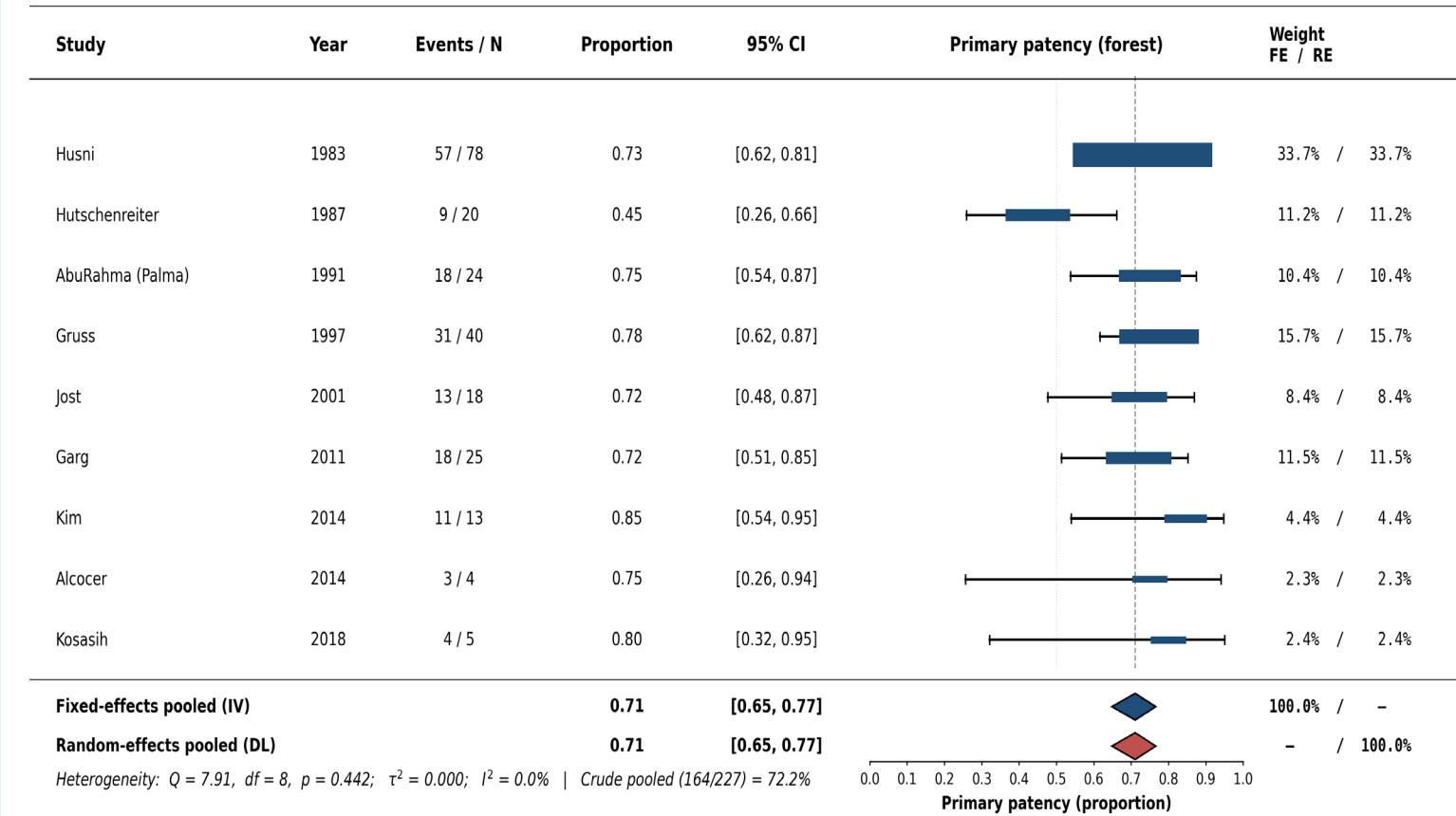
Study	Year	Center / Country	n	Patent	Patency	Conduit	Valvular repair (VVF)	NOS
Husni	1978	Cleveland, USA	78	57	73 %	GSV	Selective	6 / 9
Hutschenreiter	1979	Ulm, Germany	20	9	44 %	GSV	Routine	4 / 9
AbuRahma	1991	West Virginia, USA	24	18	75 %	GSV	None	7 / 9
Gruss	1997	Munich, Germany	40	31	78 %	GSV + PTFE	Routine	6 / 9
Jost (Mayo)	2001	Mayo Clinic, USA	18	13	72 %	GSV	Selective	9 / 9
Alcocer	2008	UAB / Mexico	4	3	75 %	GSV	None	4 / 9
Kim	2009	South Korea	13	11	84 %	GSV	Selective	7 / 9
Garg (Mayo)	2011	Mayo Clinic, USA	25	18	70 %	GSV	Selective	9 / 9
Kosasih	2017	Imperial, UK	5	4	80 %	GSV (1 PTFE)	None	6 / 9

TOTAL 227 patients · 164 patent grafts · Crude pooled patency 72.2 %

VVF = arteriovenous fistula. Routine = always, Selective = marginal inflow / small GSV, None = never. NOS: High ≥ 7 · Moderate 5-6 · Higher concern < 5

Pooled primary patency = 0.71 (95% CI 0.65–0.77)

Palma / Palma-Dale Venovenous Crossover Bypass — Primary Patency Meta-Analysis
 9 cohorts, n = 227 grafts; logit-transformed proportional meta-analysis (continuity correction 0.5)



0.71

Pooled patency (95% CI 0.65–0.77)

I² = 0.0%

No detectable between-study heterogeneity

164 / 227

Crude patency · 72.2 %

HETEROGENEITY

Q = 7.91

df = 8, p = 0.442

$\tau^2 = 0.000$ · FE = RE

Figure Logit-transformed proportional meta-analysis (continuity correction 0.5). Diamonds = pooled estimates.

Clinical benefit, complications, and mortality

75–80 %

Clinical improvement
(venous claudication, edema, ulcer healing)

≈ 28 %

Graft thrombosis (63 / 227)
Dominant failure mode

5–15 %

Wound / lymphatic complications
Serious complications uncommon

0

Palma-attributable
perioperative deaths

INTERPRETATION

Clinical improvement consistently exceeds graft patency — a single durable conduit relieves venous hypertension across most domains of post-thrombotic disease.

CONDUIT

Autologous GSV consistently outperforms PTFE.

Kosasih: only PTFE Palma in the cohort failed instantly.

Garg: prosthetic crossovers performed poorly even when used as salvage.

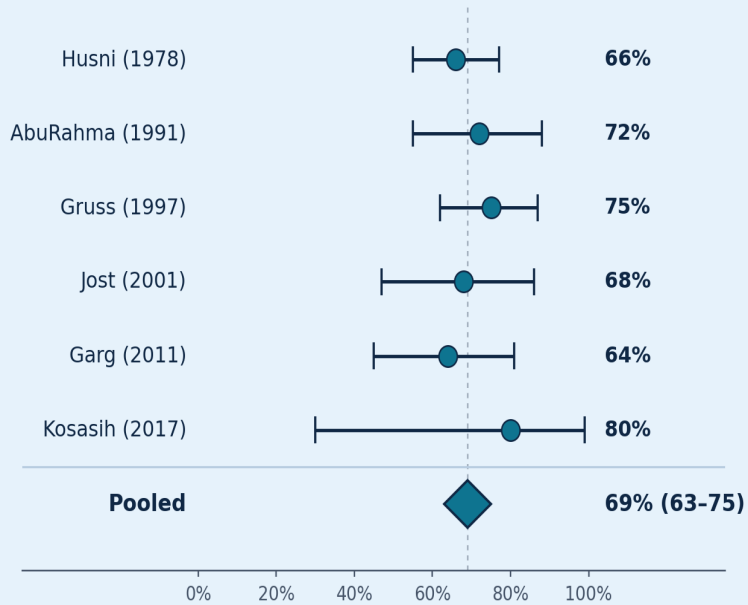
AVF POLICY

Routine AVF did NOT improve patency vs selective or no-AVF.

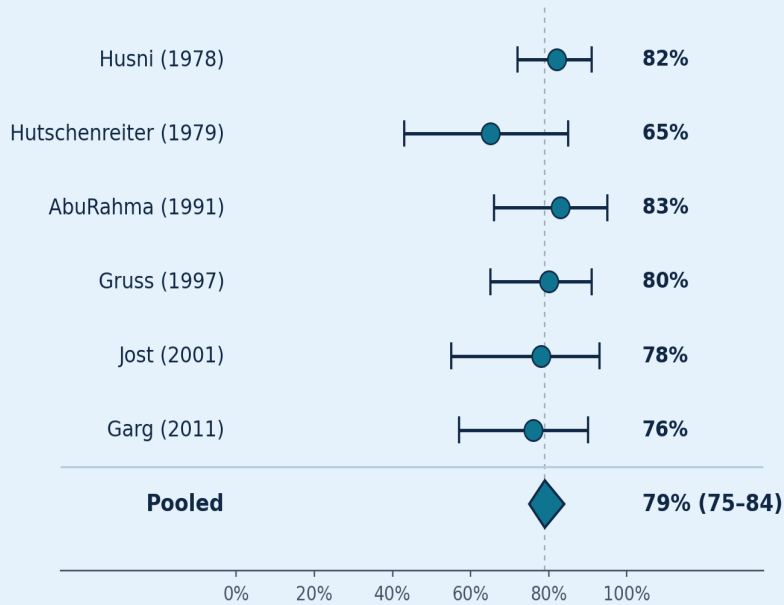
Modern technique favors selective AVF (marginal inflow / small GSV) with duplex surveillance.

Clinical outcomes – ulcer healing & edema resolution

Venous ulcer healing (patent graft)



Edema / claudication resolution



KEY POINTS

Clinical benefit tracks graft patency – healing rates drop sharply when the crossover occludes.

Ulcer recurrence \approx 50% at long-term follow-up, driven by distal reflux or graft failure.

Edema relief is the most consistently achieved endpoint in the analysis.

69 %

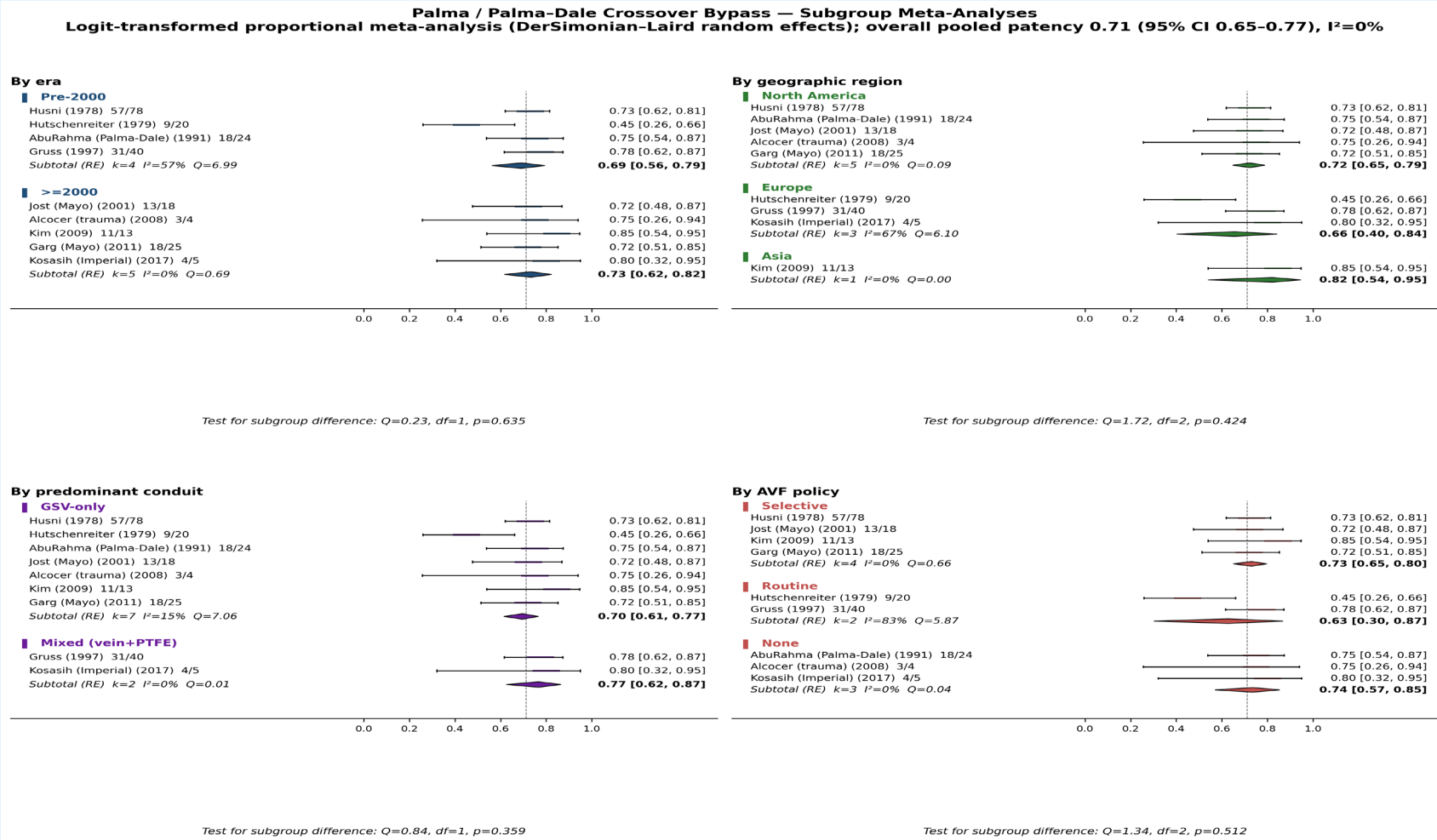
Pooled venous ulcer healing in limbs with a patent Palma graft (95 % CI 63–75)

80 %

Pooled edema / venous claudication resolution (95 % CI 75–84)

Figure . Forest plots of venous ulcer healing (patent grafts, k = 6) and edema / claudication resolution (k = 6); diamonds = DerSimonian–Laird pooled estimates.

Era · Region · Conduit · AVF policy



TEST FOR SUBGROUP DIFFERENCE

Era (pre-2000 vs ≥2000)

p = 0.635

Region (NA · EU · Asia)

p = 0.424

Conduit (GSV vs mixed)

p = 0.359

AVF policy (sel · routine · none)

p = 0.512

Take-home

Patency is consistent across decades, geography, and technique — the result is robust.

Figure . Pre-specified subgroup forest plots; vertical dashed line = pooled 0.71.

Has the modern era moved the needle? Not significantly.

+0.012 / yr

Logit slope on publication year

OR 1.12

Odds for patency per decade

p = 0.299

Not statistically significant

METHOD

Random-effects meta-regression · logit-transformed primary patency · publication year as single continuous moderator · DerSimonian–Laird τ^2 · 9 cohorts, 227 grafts

INTERPRETATION

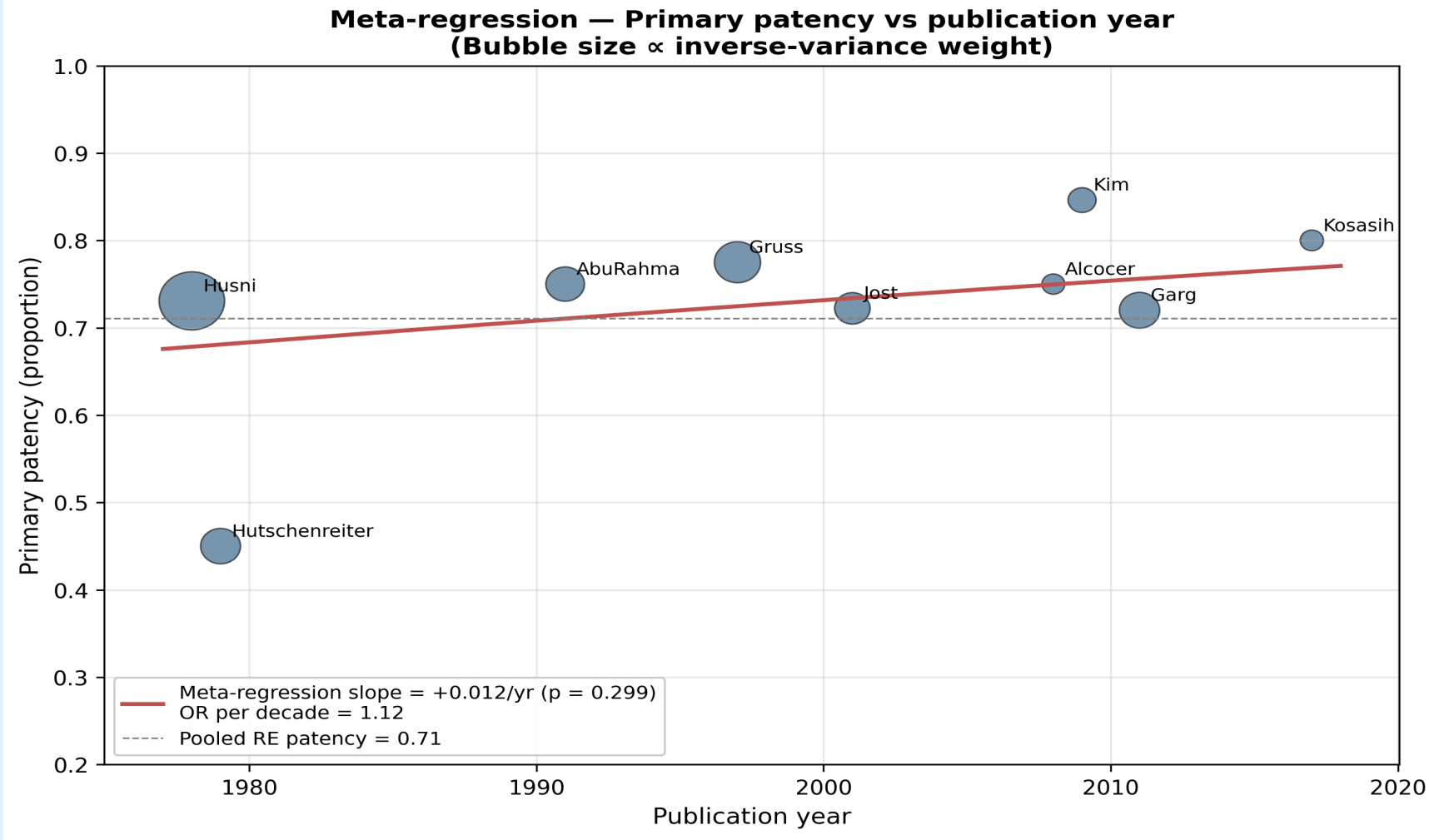
Modest positive trend, but no statistically detectable temporal change in primary patency.

Implication — patency is driven by technique and patient selection, not era.

A single durable conduit + disciplined AVF policy + autologous GSV remain the levers.

Caveat — *Hutschenreiter (1979) anchors the early-era tail and widens below-line dispersion; see Slide 11B scatter.*

Patency vs publication year – cohort-level scatter



WHAT TO NOTICE

- Gentle upward slope
- Wide 95 % CI envelopes zero
- Hutschenreiter ('79) anchors far left
- Modern cohorts cluster 60–85 % patency

TAKE-HOME

Forty-five years of iteration have not significantly changed Palma primary patency — the operation remains technique-dependent, not era-dependent.

Four scenarios where open beats endovascular

1 Long-segment chronic total occlusion

Sub-millimeter recanalization channels; unfavorable inflow for stent landing. Open conduit bypasses unstentable disease entirely.

2 Repeat in-stent occlusion or restenosis

After multiple endovascular revisions with progressive scarring, an autologous Palma is a definitive escape from the cycle.

3 Post-trauma iliac vein ligation

Damage-control ligation produces a fixed mechanical obstruction. Staged Palma-Dale (Alcocer) restores outflow once the patient is stable.

4 Anatomically unstentable / pediatric / contraindicated stenting

Hypoplastic iliac segments, cement leak after vertebroplasty, congenital agenesis, or stent contraindication.

Stent first. Reserve open reconstruction for the right anatomy.

STENT-FIRST — ENDOVASCULAR PREFERRED

- ▶ Symptomatic non-thrombotic May–Thurner
- ▶ Discrete iliofemoral post-thrombotic stenoses with viable recanalization channel
- ▶ IVC-involving or bilateral iliac disease (kissing stents, IVC reconstruction)
- ▶ Patients unfit for general anesthesia / contraindicated GSV harvest
- ▶ Recent acute DVT amenable to thrombolysis ± stenting

PALMA INAPPROPRIATE / DOOMED TO FAIL

- × Bilateral iliac obstruction (no contralateral inflow)
- × Inadequate, varicose, or absent contralateral GSV
- × IVC obstruction without iliac patency
- × Active venous infection or open contaminated field
- × When prosthetic conduit is the only option — patency is poor (Kosasih, Garg)

Our Palma reconstruction – durable at 2 years

INDICATION

74-yo male · recurrent right retroperitoneal liposarcoma · iatrogenic complete transection of the right external iliac artery and vein during oncologic resection · loss of venous outflow from the right limb.

VENOUS RECONSTRUCTION

Left-to-right cross-femoral venous bypass using contralateral great saphenous vein (Palma procedure) · end-to-side GSV–CFV anastomosis · arterial limb revascularized in the same setting · four-compartment fasciotomy.

EARLY POSTOPERATIVE COURSE

Tibial pulses triphasic; no compartment syndrome or venous congestion · fasciotomy closed at 1 week · minor wound / lymphatic issues managed conservatively · discharged home on therapeutic anticoagulation.

ONE-YEAR FOLLOW-UP

Duplex / CTA — patent Palma graft with brisk phasic flow, no significant stenosis · no recurrent limb edema, venous claudication, or ulceration · fully ambulatory and independent

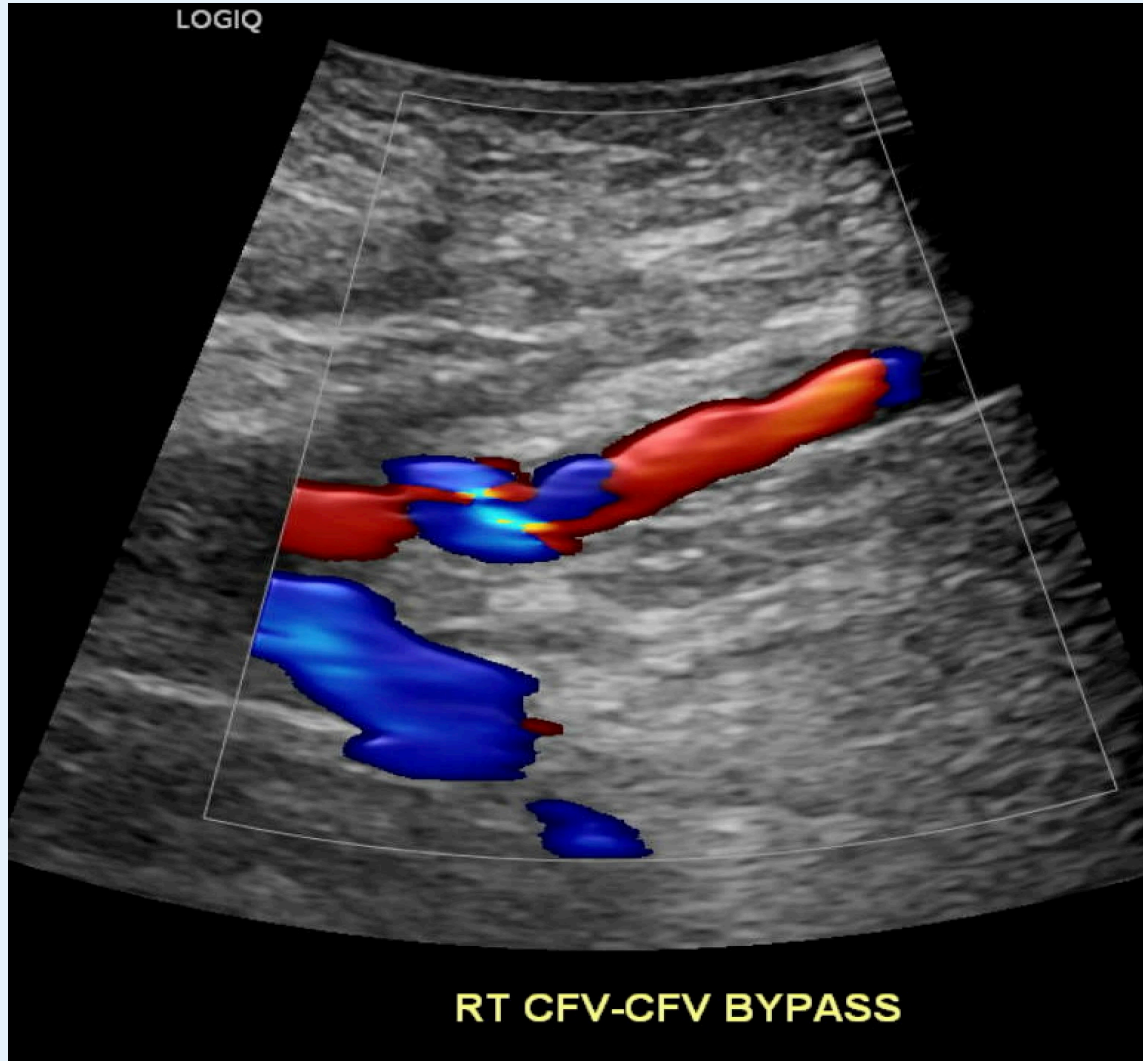


Intraoperative image:

- Left-to-right cross-femoral bypass with saphenous vein
- Successful revascularization pathway visible

).

Durable patency confirmed on follow-up imaging



brisk

IMAGING AT 12 MONTHS

- ✓ Patent cross-femoral venous bypass on duplex / CTA
- ✓ Preserved caliber · continuous phasic flow
- ✓ No graft thrombosis or hemodynamically significant stenosis
- ✓ Contralateral limb venous drainage preserved · no venous hypertension
- ✓ No limb edema, claudication, or ulceration

12 mo

Documented patent follow-up

0

Graft-related reinterventions

TAKE-HOME

Real-world confirmation from our institution — Palma can provide durable, symptom-free venous outflow in selected patients well into the endovascular era.

TAKE-HOME MESSAGES

- 1** Across 9 cohorts and 227 grafts, pooled primary patency = 0.71 (95% CI 0.65–0.77) — directly comparable to historical 70–83% benchmarks.
- 2** Clinical improvement (75–80%) consistently exceeds graft patency. Even partial outflow restoration meaningfully relieves venous hypertension.
- 3** Autologous GSV is the conduit of choice. Prosthetic Palma performs poorly and should be a salvage-of-salvage option only.
- 4** Routine arteriovenous fistula does NOT improve patency in modern data. Reserve AVF for marginal inflow or small-caliber GSV.
- 5** Stent first when feasible. But do not abandon the Palma — it remains a durable salvage when stenting is impossible, has failed, or is anatomically unsuitable.

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THANK YOU.

Questions and discussion welcome.
