

Re-Interventions after EVAR
What should we do with Type-1 Endoleaks

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Disclosure of Interest

Speaker name: Colin Bicknell

I have the following potential conflicts of interest to report:

- Medtronic: Consultancy, Speakers fees, travel and conference fees
- Orzone: Institutional level capital funding
- Bolton Medical: Consultancy, speakers fees, travel and conference fees
- Gore: Travel and conference fees



Case presentation - 2009

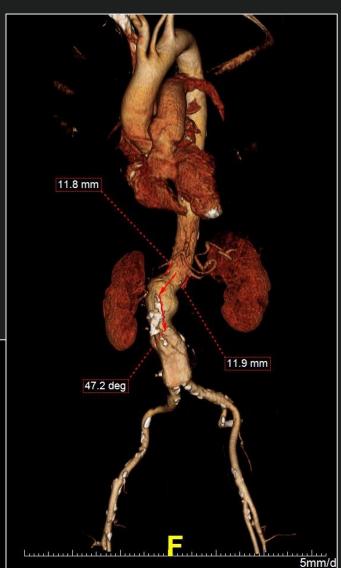
69 year old man

Hypertension, High Cholesterol, Ex smoker, Asthma

- Creat 129 mmol/L
- No inducible ischaemia on DSE
- FEV₁ 1.5L, FEV₁ / FVC ratio 65%



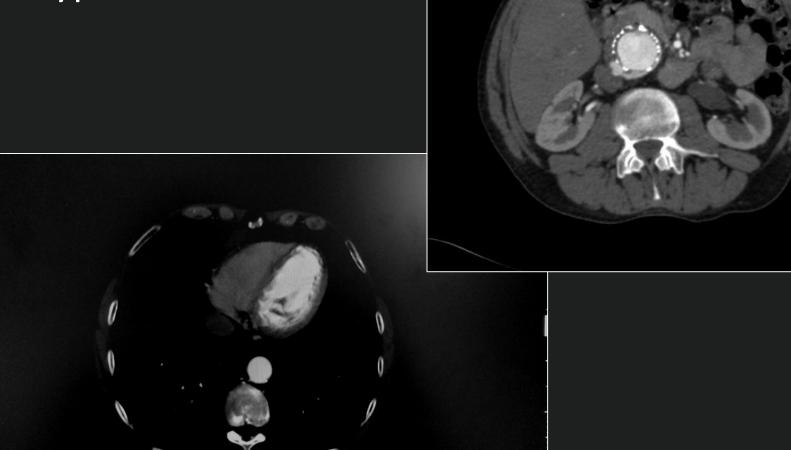






2010

• Type 1 Endoleak

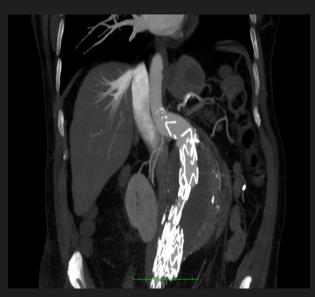


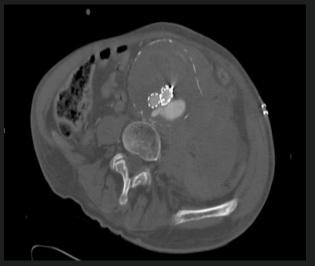


ENDOLEAK TYPES

...not all endoleaks are the same

- Endograft mal-deployment
- Excessive thrombus/calcium in seal zone
- Inadvertent creation of a leak channel
 - Excessive oversizing creating gutters
 - Non-circular or angulated neck conformability challenges
- Migration and loss of seal
- Late insufficient apposition due to aortic expansion







What do we do with early type 1 endoleaks after EVAR?



What do we do with early type 1 endoleaks after EVAR?

FIRST:

Correct Endograft Mal-deployment

- Re-balloon
- Extend to renals ensuring max seal
- Palmaz



Balloon, extend, Palmaz









SECONDLY, FOR LEAK CHANNEL

HELI-FX™ ENDOANCHOR™ IMPLANT SYSTEM ENDOVASCULAR INTERRUPTED SUTURE SYSTEM





Principles of Endoleak treatment

Spatial approximation of source



Targeted EndoAnchor placement



Circumferential reinforcement

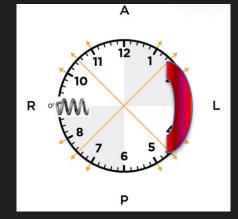


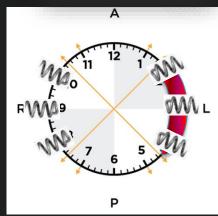
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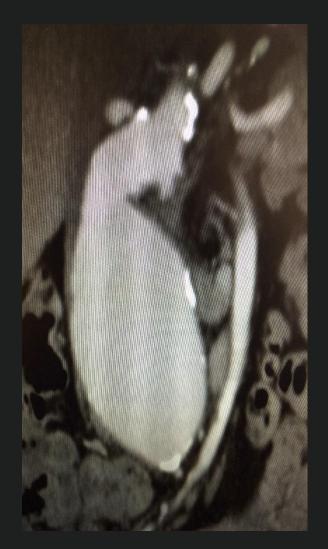
Approximated endoleak location







CONFORMABILITY CHALLENGES

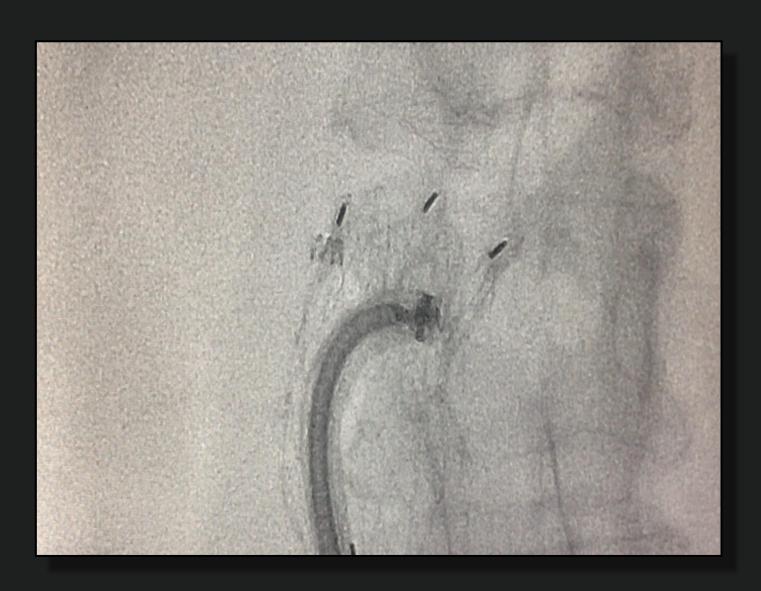






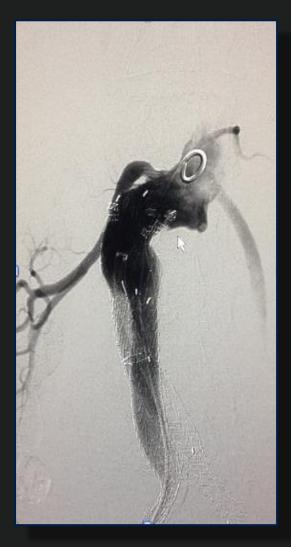


CONFORMABILITY CHALLENGES





CONFORMABILITY CHALLENGES

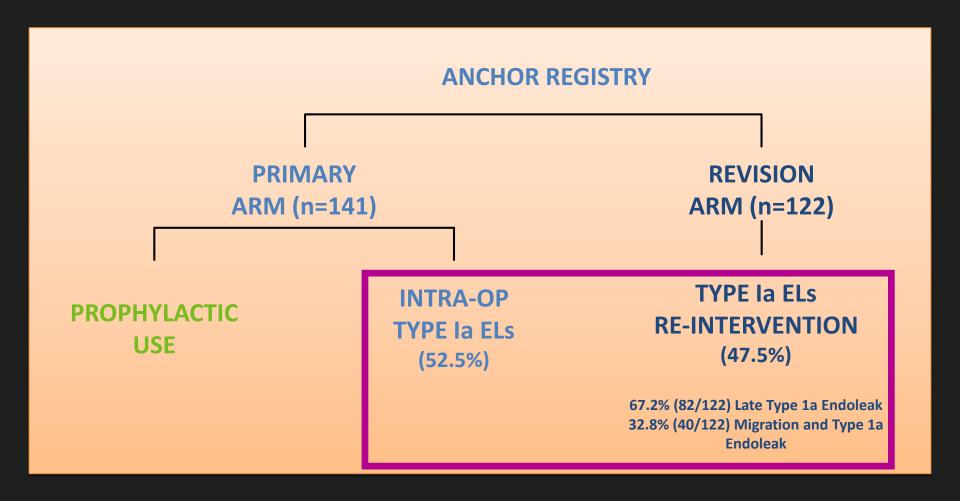






ANCHOR Registry – Therapeutic USE

Therapeutic Use cohort (N=263) consists of patients receiving EndoAnchor™ implants to treat Type 1a Endoleaks in the Primary and Revision Arms.





ANCHOR Registry – Therapeutic Use for Proximal ELs

TECHNICAL SUCCESS

Deployment of desired number of EndoAnchor™ implants without fracture or loss of integrity

95.7% Intra-op T1 EL

93.4% Revision

PROCEDURAL SUCCESS

Technical success without type Ia endoleak at completion arteriography

85.1% Intra-op T1 EL

82.8% Revision

Avg. duration of Procedure (min)

159 158

Intra-op Revision

T1 EL Site-reported data Avg. time to EndoAnchor™ implants (min)

T1 EL

20 23
Intra-op Revision

Avg. number of EndoAnchor™ implants

6.1 7.7

Intra-op Revision T1 EL



ANCHOR Registry – Therapeutic Use for Proximal ELs

Persistent/Recurrent Type Ia Endoleaks

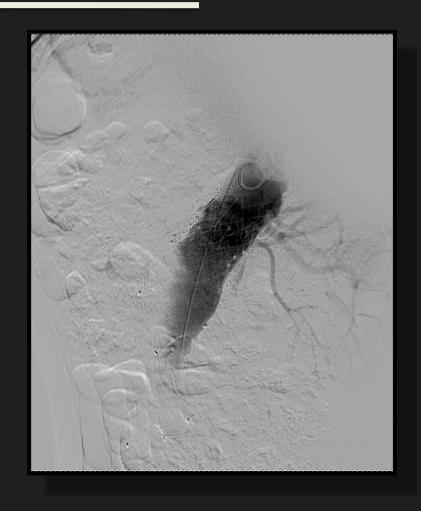
Mean follow-up 10.4 months

	Type 1a ELs	CTs	%
Intra-op T1 EL	3	76	3.9%
Revision	21	66	31.8%
All	24	142	16.9%

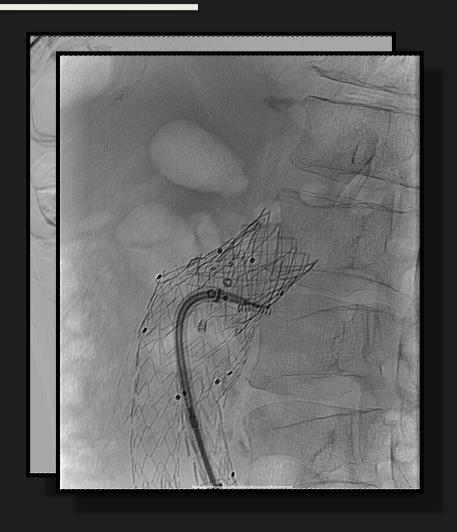
18/24 had no subsequent interventions to repair the persistent/recurrent type

1a endoleaks

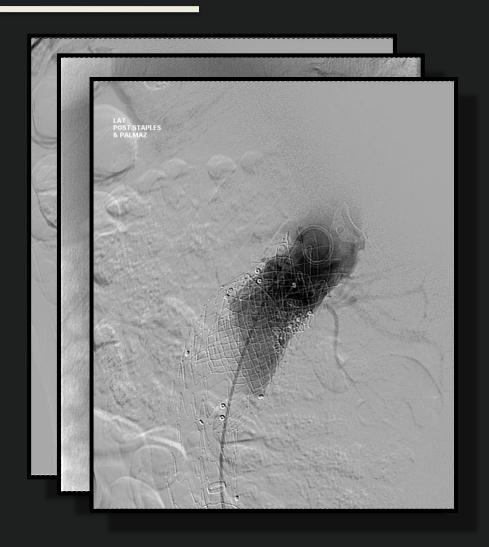
ENDOANACHOR USE: REPAIR



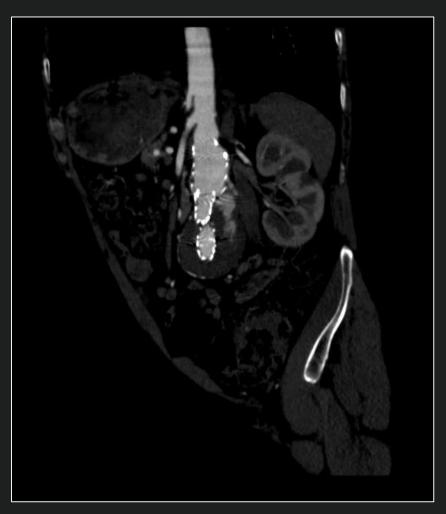
ENDOANACHOR USE: REPAIR

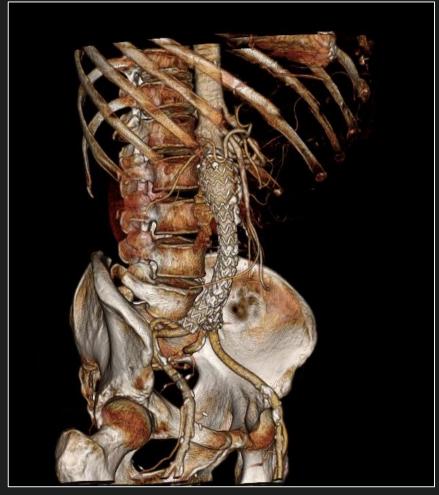


ENDOANACHOR USE: REPAIR



Further Sac Expansion by 2011 with persistent type 1 endoleak







LATE type 1

- Migration and loss of seal
 - Extend, re- balloon
- Neck degeneration
 - Fenestrated cuff
 - Chimney / CHEVAS
 - Hybrid
 - Open explant
 - Embolisation





Fenestrated Cuff



- Imperial Experience
- 2010-16
- Type IA endoleak following EVAR
- 10 patients
- Mean age 78 years; seven patients
- ASA grade ≥ III
- Five cuffs, five fenestrated re-lining
- Technical success was 9/10
- Median hospital stay of 6.5 (6-16) days
- No 30-day mortality
- Mean follow up was 22.4 ± 13 months.
- One patient died at 51 months following the FEVAR from a ruptured abdominal aortic aneurysm. Known to have type 2 endoleak
- No other aneurysm related death



The Chimney Option

Techniques evolving

Experience increasing

Results more convincing

Not in this case - Palmaz





PERICLES

Collected World Experience About the Performance of the Snorkel/Chimney Endovascular Technique in the Treatment of Complex Aortic Pathologies

The PERICLES Registry

Konstantinos P. Donas, MD, Jason T. Lee, MD, † Mario Lachat, MD, ‡ Giovanni Torsello, MD, PhD, § and Frank J. Veith, MD: on behalf of the PERICLES investigators

Konstantinos P. Donas, MD.* Jason T. Lee, MD.† Mario Lachat, MD.‡ Giovanni Torsello, MD, PhD, S and Frunk I. Veith MD.* on beholf of the DEPLICE Desirement

- Retrospective
- 517 patients treated by ch-EVAR from 2008-2014
- Mean follow-up of 17.1 months (range: 1-70 months)
- Primary patency 94%, secondary patency 95.3%.

Results affected by:

- Conformity technique
- Device sizing
- Device selection

Intra-op type la endoleak:	7.9%
Persistent intra-op type la endoleak:	2.9%
Type IA endoleak at latest FU:	5.8%
Technical Success	97%

...for 517 patients from 13 international centers





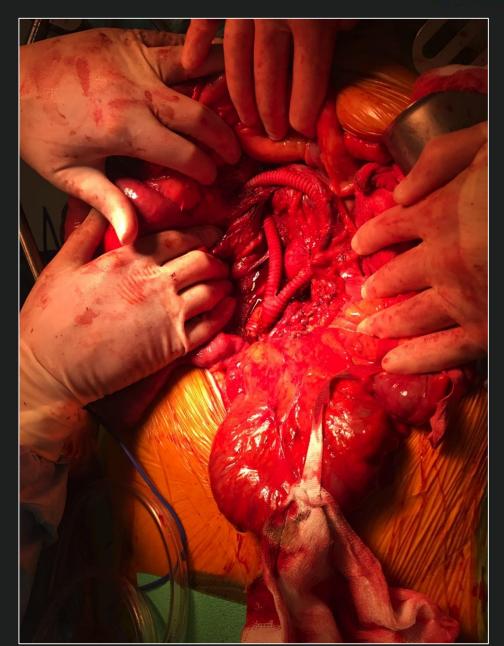






The hybrid approach

- Retrograde/antegrade or extra-anatomical revascularisation of renals, SMA and/or coeliac
- Stenting to normal segment
- An adequate rescue, careful planning needed





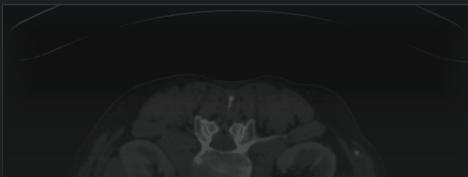
The conclusion...



CT 2012

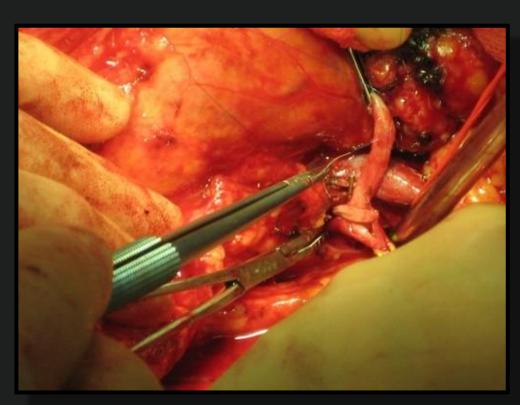


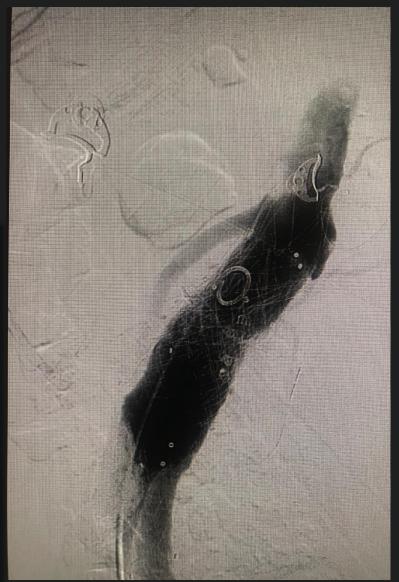
Neck dilated Renal artery disease (5mm) Now 73 Years, and less fit!



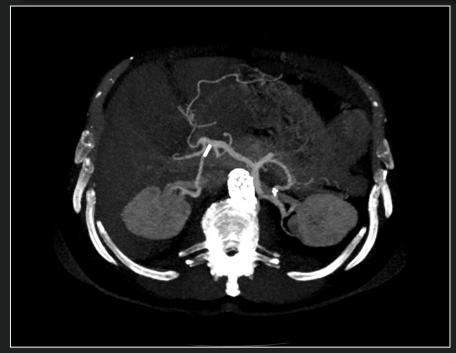


Hepato/spleno – renal bypass and stenting













Previous EVAR and Type 1 Endoleak

Early/intra-operative

Late, due to migration

Late, due to degeneration

Re-balloon

Extension piece

Extend

Endoanchor if conformability issues or leak channel identified Algorithm for type 1

Fenestrated cuff

Chimney

Hybrid or open approach

Fill with onyx if no other solution