



Australian Government
Department of Health and Ageing



Australia and New Zealand Horizon Scanning Network

ANZHSN

AN INITIATIVE OF THE NATIONAL STATE AND
TERRITORY GOVERNMENTS OF AUSTRALIA
AND THE GOVERNMENT OF NEW ZEALAND

Horizon Scanning Technology Prioritising Summaries

Intracranial angioplasty and stenting
(WingSpan™ self-expanding stent) for
cerebral atherosclerotic stenosis

June 2006



ASERNIP(S)

**Australian
Safety
and Efficacy
Register
of New
Interventional
Procedures -
Surgical**



**Royal Australasian
College of Surgeons**



© Commonwealth of Australia [2006]

This work is copyright. You may download, display, print and reproduce this material in unaltered form only (retaining this notice) for your personal, non-commercial use or use within your organisation. Apart from any use as permitted under the Copyright Act 1968, all other rights are reserved. Requests and inquiries concerning reproduction and rights should be addressed to Commonwealth Copyright Administration, Attorney General's Department, Robert Garran Offices, National Circuit, Canberra ACT 2600 or posted at <http://www.ag.gov.au/cca>

Electronic copies can be obtained from <http://www.horizonscanning.gov.au>

Enquiries about the content of the report should be directed to:

HealthPACT Secretariat
Department of Health and Ageing
MDP 106
GPO Box 9848
Canberra ACT 2606
AUSTRALIA

DISCLAIMER: This report is based on information available at the time of research and cannot be expected to cover any developments arising from subsequent improvements to health technologies. This report is based on a limited literature search and is not a definitive statement on the safety, effectiveness or cost-effectiveness of the health technology covered.

The Commonwealth does not guarantee the accuracy, currency or completeness of the information in this report. This report is not intended to be used as medical advice and it is not intended to be used to diagnose, treat, cure or prevent any disease, nor should it be used for therapeutic purposes or as a substitute for a health professional's advice. The Commonwealth does not accept any liability for any injury, loss or damage incurred by use of or reliance on the information.

The production of this Horizon scanning prioritising summary was overseen by the Health Policy Advisory Committee on Technology (HealthPACT), a sub-committee of the Medical Services Advisory Committee (MSAC). HealthPACT comprises representatives from health departments in all states and territories, the Australia and New Zealand governments; MSAC and ASERNIP-S. The Australian Health Ministers' Advisory Council (AHMAC) supports HealthPACT through funding.

This Horizon scanning prioritising summary was prepared by staff from the Australian Safety and Efficacy Register of New Interventional Procedures – Surgical (ASERNIP-S).

Name of Technology:



Intracranial angioplasty and stenting (WingSpan™ self-expanding stent) for cerebral atherosclerotic stenosis.

Purpose and Target Group:

Patients suffering from symptomatic intracranial atherosclerotic stenosis.

Stage of Development (in Australia):

- Experimental
- Investigational
- Nearly established
- Established
- Established but changed indication or modification of technique
- Should be taken out of use
- Not yet emerged

The WingSpan stent is currently not available in Australia. Hence it is not listed or registered in the Australian Register of Therapeutic Goods database.

International Utilisation:

COUNTRY	LEVEL OF USE		
	Trials underway	Limited use	Widely diffused
United States	✓		
Germany	✓		

Impact Summary:

Background

Intracranial cerebral atherosclerosis is a disease that is characterised by endothelial dysfunction, vascular inflammation, and the build-up of lipids, cholesterol, calcium and cellular debris within the intima of the vessel wall (Orford 2005). It is estimated that intracranial cerebral atherosclerosis accounts for approximately 8% to 10% of all ischemic strokes, with a higher incidence in Asian, African and Hispanic populations (Higashida *et al.* 2005). The aetiology of ischemic strokes secondary to intracranial atherosclerosis have presented four potential mechanisms, namely hypoperfusion, thrombosis at the intraplaque haemorrhage or occlusive plaque growth, thromboembolic events distal to the site of stenosis or direct occlusion of small penetrating arteries at the site of the plaque (Higashida *et al.* 2005).

Research has revealed that despite treatment, patients with symptomatic intracranial atherosclerosis have unacceptably high rates of recurrent cerebrovascular ischemic events, coronary heart disease and death (Leung *et al.* 2006). To date, the best medical treatment for intracranial atherosclerosis remains controversial and aspirin is commonly used as the standard treatment. Anticoagulative medications have shown little success at reducing



recurrent vascular events, the Warfarin-Aspirin Symptomatic Intracranial Disease (WASID) Trial failed to show a significant risk reduction of primary end-points (stroke and vascular death) when comparing warfarin treatment to aspirin. In addition, the WASID trial was prematurely terminated due to the mortality rates of the warfarin group (9.7%, 28/289) and the high incidence of major bleeding complications (8.3%, 24/289) compared to the aspirin group (Leung *et al.* 2006).

Due to poor patient outcomes with medical therapy, endovascular treatments have been proposed as an effective alternative. Percutaneous transluminal angioplasty (balloon angioplasty) was developed as a means of increasing lumen size at the region of stenosis via compression of the plaque. A recent study by Marks *et al.* (2005) found that percutaneous transluminal angioplasty is capable of reducing annual stroke rates to 3.36%, a substantial reduction compared to the approximate annual stroke rate for intracranial stenosis of 8% to 12% (Kirmani *et al.* 2005). However, restenosis rates for this technique ranges from 30% to 50% (Leung *et al.* 2006) and despite encouraging results, it is a hazardous procedure with reports of up to 50% complication rates and 17% fatality rates (Gupta *et al.* 2003).

An alternate technique involves the insertion of a stent at the site of the lesion; however this method is often technically difficult due to the limited flexibility of these stents. The introduction of more flexible stents (e.g. NeuroLink) designed specifically for intracranial use has enabled safer insertion and therefore treatment of intracranial atherosclerosis. A clinical trial utilising the NeuroLink (Guidant, CA) has a documented 95% success rates and one month post-operative stroke rates of 7%, as shown in the Stenting in Symptomatic Atherosclerotic Lesions of Vertebral Intracranial Arteries (SSYLVIA) study (The SSYLVIA study investigators 2004). However, restenosis rates remain unsatisfactory with 32% of patients developing $\geq 50\%$ restenosis after 6 months (Henkes *et al.* 2005).

A new concept for percutaneous intracranial angioplasty involves the use of a self-expanding stent. This new stent, the WingSpan (Boston Scientific, Fremont, CA), is a self-expanding, neurovascular, flexible nitinol stent with purportedly excellent trackability in intracranial vasculature (Leung *et al.* 2006). This procedure begins with a stentless angioplasty, followed by covering of the previously dilated stenosis with the WingSpan stent (Henkes *et al.* 2005). The rationale for this new procedure is that the radial self-expanding force which the stent exerts after deployment would result in a further gradual reduction in stenosis, therefore a smaller balloon can be used during the pre-dilation stage. The use of a smaller balloon should reduce the risk of vascular trauma, a state where the balloon can cause intimal dissection, recoiling or vessel rupture, which has been a recurring disadvantage in standard balloon angioplasty (Leung *et al.* 2006).

Clinical Need and Burden of Disease

Stroke is the third leading cause of death and the leading cause of adult disability in North America, Europe and Asia (Higashida *et al.* 2005). As previously stated, intracranial cerebral



atherosclerosis accounts for approximately 8% to 10% of all ischemic strokes. This translates to 40,000 to 60,000 new stroke cases annually in the United States (Higashida *et al.* 2005). In Australia, the 2001 National Health Survey revealed that approximately 1.2% of Australians have suffered a stroke, this corresponds to 217, 500 Australians. There are approximately 40,000 to 48,000 stroke events a year in Australia (AIHW 2006); therefore approximately 3200 to 3840 of these are caused by intracranial atherosclerosis. Studies have shown that the annual stroke risk from all causes in patients with intracranial atherosclerosis ranges from at least 3.6% to over 13% annually (Higashida *et al.* 2005).

Estimated Speed, Geographic and Practitioner Use, Patterns of Diffusion in the Health System

The WingSpan stent is currently in the experimental stages and therefore has not been approved for marketing in any country. However, it has received humanitarian device exemption approval (HDE number: H050001) from the FDA in 2004 (Food and Drug Administration 2006).

The WingSpan stent is the only stent currently available which is designed specifically for the treatment of intracranial stenosis in conjunction with balloon angioplasty. If proven safe and effective, it may provide a safer alternative to standard balloon angioplasty or stenting.

Existing Comparators

Treatment of recurring ischemic stroke resulting from intracranial atherosclerosis currently includes:

- Medical therapy (antiplatelets and anticoagulants)
- Percutaneous transluminal angioplasty
- Coronary stent insertion
- Surgical bypass of the stenosis

Estimated Cost Impact

A representative from the University of Wisconsin Hospital stated that the total cost for the entire procedure (balloon angioplasty and WingSpan deployment) ranges from US\$48,000 to US\$ 125,000. The WingSpan stent itself costs US\$5000 (Kansas City Star 2006).

The Medicare Benefits Schedule does not list any reimbursements for the use of stents or balloon angioplasty for intracranial atherosclerosis. However, the reimbursement fees for extracranial to intracranial bypass using a superficial temporal artery (Item number: 39818) is AU\$1581.45 while the reimbursements for extracranial to intracranial bypass using a saphenous vein graft (Item number: 39821) is AU\$1877.85. From July 2004 to June 2005, there were 14 Medicare claims for extracranial to intracranial bypass using a superficial



temporal artery and 5 Medicare claims for extracranial to intracranial bypass using a saphenous vein graft (Medicare Australia 2006).

Efficacy and Safety Issues

List of Studies Found

Total number of studies	2
Case series	2

The studies included in this summary are highlighted in bold in the reference list.

Safety and efficacy data from 2 case series studies have been selected for inclusion in this summary.

A 45 patient multicentre clinical trial was conducted to determine the safety and efficacy of intracranial angioplasty and stenting utilising the Gateway™ PTA Balloon catheter and the WingSpan stent. The results of this trial was not published in any peer reviewed journals, instead data was extracted from the FDA Safety and Efficacy Summary. The investigators did not include a control group for this trial due to the fact that there is no alternative standard therapy for this disease, results were compared to historical controls extracted from peer-reviewed literature with similar patient cohorts. Of the 45 patients, intracranial angioplasty and stenting was feasible in 44 patients (97.8%). One patient was deemed unsuitable for the treatment due to tortuous anatomy. At 30 days post-treatment, 2/44 (4.5%) patients died due to ipsilateral stroke (stroke occurred at the same hemisphere of the targeted lesion), another 2 patients (4.5%) experienced major ipsilateral stroke, and there was one death (2.2%) due to cerebral haemorrhage 10 days post-treatment. At 6 months post-treatment, 42 patients were evaluated and the overall stroke rate was 9.5% (4/42 patients) with one death (2.4%). Analysis of vessel characteristics at 6 month post-treatment revealed the following results (Table 2) (Food and Drug Administration 2006):

Table 2: Vessel characteristics up to 6 months post-treatment



	Baseline (n=45)	Post PTA (n=44)	Post intracranial stenting (WingSpan) (n=44)	6 months post- treatment (n=40)
Reference vessel diameter (mm)	3.1 ± 0.8	3.2 ± 0.8	3.2 ± 0.8	3.1 ± 0.8
MLD at target lesion (mm)	0.8 ± 0.6	1.6 ± 0.6	2.1 ± 0.5	2.2 ± 0.8
% Stenosis	74.9 ± 9.8	50.0 ± 16.2	31.9 ± 13.6	28.0 ± 23.2
≥ 50% stenosis	100% (45/45)	54.5% (24/44)	0.0% (0/44)	7.5% (3/40)

The results indicate that there was a slight increase in MLD and a slight decrease in percentage stenosis at 6 months post-treatment; however these results were not significant. Despite this, the WingSpan stent was successful in maintaining $\leq 50\%$ stenosis in 92.5% (37/40 patients) of patients 6 months post-treatment (Food and Drug Administration 2006). In comparison to the SSYLVIA study (normal stenting without prior balloon angioplasty), the overall stroke and death rate was lower (Table 3) but these results must be evaluated in the light of the shorter follow-up duration of the current study (174 days for WingSpan versus 216 days for SSYLVIA).

Table 3: SSYLVIA study vs Wingspan (45 patient) clinical trial

	All stroke rate	Death rate	All stroke and death	Ipsilateral stroke
SSYLVIA study (n=61)	13.1% (8/61)	6.6% (4/61)	13.1% (8/61)	11.5% (7/61)
WingSpan (n=45)	9.5% (4/42)	2.4% (1/42)	9.5% (4/42)	7.1% 3/42)

No parent vessel dissections or stent migration was reported during the WingSpan 45 patient trial. Four cases (4/44, 9%) of access site related complications were encountered and all required treatment. In addition, five patients (11%) suffered 7 access site related adverse events with four requiring treatment (Food and Drug Administration 2006).

The case series by Henkes *et al.* (2004) reported that balloon angioplasty and WingSpan stenting reduced stenosis from a baseline of 72% to 54% after balloon dilatation and to 38% after stent deployment in 15 patients. One patient who had a middle cerebral artery (MCA) stenosis suffered MCA branch occlusion as a result of the procedure, resulting in a transient increase of pre-existing hemiparesis. Ischemic symptoms that were observed in the remaining 14 patients prior to treatment were completely resolved after balloon angioplasty and stenting. No additional follow-up results were presented in this study. The authors



reported that the procedure went smoothly with a 100% success rate and no incidence of dissection or elastic recoil. Deployment of the WingSpan stent was uneventful, with no visible damage to the vessel and no cases of in-stent thrombosis (Henkes *et al.* 2004).

Ethical Issues

No issues were identified from the retrieved materials.

Cultural or Religious Considerations

No issues were identified from the retrieved materials.

Other Issues

No issues were identified from the retrieved materials.

Recommendation

Balloon angioplasty and WingSpan stent deployment offers a potentially safe treatment for intracranial atherosclerotic stenosis. However, the evidence available is limited and despite good clinical outcomes there is a need for larger multicentre trials with long-term follow-up to determine long-term patency and stroke rates. Comparative studies with standard balloon angioplasty or stenting would be valuable in determining the value of this procedure as well. Due to the potential benefits of this procedure, it is recommended that it is monitored for 12 months.

- | | |
|--|--|
| <input type="checkbox"/> Horizon Scanning Report | <input type="checkbox"/> Full Health Technology Assessment |
| <input checked="" type="checkbox"/> Monitor | <input type="checkbox"/> Archive |

References:

AIHW (Australian Institute of Health and Welfare). Incidence and prevalence of chronic diseases. Last updated 2005.

http://www.aihw.gov.au/cdarf/data_pages/incidence_prevalence/index.cfm#Stroke
[Accessed April 2006].

Food and Drug Administration. WingSpan™ Stent System with Gateway™ PTA Balloon Catheter. Summary of safety and probable benefit. Last updated 2006.

<http://www.fda.gov/cdrh/pdf5/h050001b.pdf> [Accessed April 2006].



Gupta R, Schumacher HC, Mangla S, Meyers PM, Duong H, Khandji AG, Marshall RS, Mohr JP, Pile-Spellman J. Urgent endovascular revascularization for symptomatic intracranial atherosclerotic stenosis. *Neurology* 2003;61: 1729-1735.

Henkes H, Miloslavski E, Lowens S, Reinartz J, Liebig T, Kuhne D. Treatment of intracranial atherosclerotic stenoses with balloon dilatation and self-expanding stent deployment (WingSpan). *Neuroradiology* 2005;47:222-228.

Higashida RT, Meyers PM, Connors JJ, Sacks D, Strother CM, Barr JD, Wojak JC, Duckwiler GR. Intracranial angioplasty and stenting for cerebral atherosclerosis: A position statement of the American Society of Interventional and Therapeutic Neuroradiology, Society of Interventional Radiology, and the American Society of Neuroradiology. *Journal of Vascular and Interventional Radiology* 2005;16(10): 1281-1285.

Kansas City Star. New stent intended to increase blood flow, reduce stroke risk. Last updated 2006. <http://www.kansascity.com/mld/kansascity/13896399.htm> [Accessed April 2006].

Kirmani JF, Janjua N, Kawi AA, Ahmed S, Khatri I, Ebrahimi A, Divani AA, Qureshi AI. Therapeutic advances in interventional neurology. *NeuroRx*® 2005;2(2): 304-323.

Leung TW, Kwon SU, Wong KS. Management of patient with symptomatic intracranial atherosclerosis. *International Journal of Stroke* 2006;1(1): 20-25.

Marks MP, Marcellus ML, Do HM, Schraedley-Desmond PK, Steinberg GK, Tong DC, Albers GW. Intracranial angioplasty without stenting for symptomatic atherosclerotic stenosis: Long-term follow-up. *American Journal of Neuroradiology* 2005;26: 525-530.

Medicare Australia. Last updated 2006. <http://www9.health.gov.au/mbs/> [Accessed April 2006].

The SSYLVA study investigators. Stenting of symptomatic atherosclerotic lesion in the vertebral or intracranial arteries (SSYLVA). Study Results. *Stroke* 2004;35: 1388-1392.

Orford JL. Atherosclerosis. Last updated 2005.
<http://www.emedicine.com/med/topic182.htm> [Accessed April 2006].



Search Criteria:

A search of MEDLINE, PubMed, *The Cochrane Library*, the Current Controlled Trials metaRegister, the UK National Research Register, the International Network of Agencies for Health Technology Assessment, relevant online journals and the Internet was conducted in April 2006.

Search terms used were: 'balloon angioplasty and stent', 'WingSpan stent', 'self-expanding intracranial stent', 'intracranial stent', 'intracranial angioplasty and stent', 'balloon dilatation and stent'.

This Horizon Scanning Prioritising Summary was prepared by Mr. Irving Lee from the NET-S Project, ASERNIP-S for the Health Policy Advisory Committee on Technology (Health PACT), on behalf of the Medical Services Advisory Committee (MSAC) and the Australian Health Ministers' Advisory Council (AHMAC).