



Deposited via The University of Leeds.

White Rose Research Online URL for this paper:

<https://eprints.whiterose.ac.uk/id/eprint/84181/>

Version: Accepted Version

---

**Article:**

Killan, CF, Killan, EC and Raine, CH (2015) Changes in children's speech discrimination and spatial release from masking between 2 and 4 years after sequential cochlear implantation. *Cochlear Implants International*, 16 (5). pp. 270-276. ISSN: 1467-0100

<https://doi.org/10.1179/1754762815Y.0000000001>

---

**Reuse**

Items deposited in White Rose Research Online are protected by copyright, with all rights reserved unless indicated otherwise. They may be downloaded and/or printed for private study, or other acts as permitted by national copyright laws. The publisher or other rights holders may allow further reproduction and re-use of the full text version. This is indicated by the licence information on the White Rose Research Online record for the item.

**Takedown**

If you consider content in White Rose Research Online to be in breach of UK law, please notify us by emailing [eprints@whiterose.ac.uk](mailto:eprints@whiterose.ac.uk) including the URL of the record and the reason for the withdrawal request.



**UNIVERSITY OF LEEDS**



**University of  
Sheffield**



**UNIVERSITY  
of York**

1   **Changes in children's speech discrimination and spatial release from masking between**  
2   **two and four years after sequential cochlear implantation**

3

4   Catherine F Killan<sup>a</sup>, Edward C Killan<sup>b</sup> and Christopher H Raine<sup>a</sup>

5

6   <sup>a</sup>Yorkshire Auditory Implant Service, Bradford Royal Infirmary, UK

7   <sup>b</sup>Faculty of Medicine and Health, University of Leeds, UK

8

9

10   Correspondence to: Catherine Killan, Yorkshire Auditory Implant Service, Listening for Life  
11   Centre, Bradford Royal Infirmary, Duckworth Lane, Bradford, West Yorkshire, BD9 6RJ,  
12   UK.

13   Tel: +44 0 1274 364853

14   Email: catherine.killan@bthft.nhs.uk

15

16   No conflicts of interest to declare.

17

18   Funding: The Ear Trust, Registered Charity No. 1000929

19

20   Acknowledgement: We thank Nicola Royle for her contribution to spatial listening  
21   assessment at the Yorkshire Auditory Implant Service.

22

## 23 ABSTRACT

24

25 Objective: To document changes in speech reception thresholds (SRTs) and spatial release  
26 from masking (SRM) for sequentially implanted children at two and four years after they  
27 received their second cochlear implant (CI<sub>2</sub>).

28 Methods: Participants were 17 children who consistently used two sequentially implanted and  
29 optimally programmed cochlear implants. SRTs were measured monaurally in quiet and  
30 binaurally in noise using the adaptive McCormick Toy Discrimination Test. Speech signals  
31 were presented from 0° azimuth and noise from 0°, +90° or -90° azimuth. SRM was  
32 calculated from SRTs in noise. Measurements were made at two and four years post-Cl<sub>2</sub>.

33 Results: There were significant improvements over time in SRTs in quiet, SRTs in noise and  
34 SRM. SRTs in quiet improved more for Cl<sub>2</sub> than for the first implant (Cl<sub>1</sub>). SRTs in noise and  
35 SRM improved more when noise was presented closest to Cl<sub>1</sub> than when closest to Cl<sub>2</sub>.  
36 Performance became more symmetrical over time.

37 Discussion: Despite prolonged periods of unilateral auditory deprivation sequentially-  
38 implanted children exhibited continued improvement in SRT and SRM. These results are  
39 valuable in setting expectations for and counselling families of children considering  
40 sequential cochlear implants.

41

42 Keywords: Cochlear Implants; Bilateral; Spatial Release from Masking; Speech  
43 Discrimination; Sequential; Speech Reception Thresholds; Speech Intelligibility

44

## 45 INTRODUCTION

46

47 One advantage of binaural hearing is an increased ability to discriminate speech from  
48 background noise due to spatial release from masking (SRM). SRM refers to the  
49 improvement in speech discrimination obtained when speech and noise signals are spatially  
50 separated, and has been attributed to the head-shadow effect and binaural processing (e.g.  
51 Hawley *et al.*, 2004; Akeroyd, 2006). One aim of bilateral cochlear implantation in children  
52 is to realize this benefit for profoundly deaf children. Bilateral cochlear implantation can be  
53 performed simultaneously but is often performed sequentially (i.e. implantation occurs one  
54 ear at a time, with the second implant, CI<sub>2</sub>, being implanted some time, often years, following  
55 the first, CI<sub>1</sub>). As a result, sequentially-implanted children may experience prolonged and  
56 asymmetrical auditory deprivation compared to normally-hearing children, children who use  
57 bilateral hearing aids and children who undergo simultaneous cochlear implantation. As a  
58 consequence, the development of binaural listening skills for sequentially-implanted children  
59 is more likely to be limited by changes in plasticity in the maturing auditory system (Sharma  
60 *et al.*, 2007; Green *et al.*, 2011; Gordon *et al.*, 2013; Sparreboom, 2013).

61

62 Several studies have described changes in speech discrimination for sequentially-implanted  
63 children as a function of time up to two years post-CI<sub>2</sub> (Peters *et al.*, 2007; Sparreboom *et al.*,  
64 2011; Strom-Roum *et al.*, 2012). In general, these studies show improvements in monaural  
65 and binaural speech reception thresholds (SRTs) in quiet and noise. Further, whilst children  
66 tend to perform better when listening via CI<sub>1</sub> alone compared to via CI<sub>2</sub> alone, the greatest  
67 improvements over time are seen for children listening via CI<sub>2</sub>. To date, longitudinal data  
68 describing speech discrimination over a time period longer than two years post-CI<sub>2</sub> have not  
69 been reported in the literature. Even less is known regarding the development over time of

70 SRM for sequentially implanted children. A number of studies have shown that sequentially  
71 implanted children display asymmetrical SRM, i.e. greater SRM is available when the noise  
72 signal is closer to CI<sub>2</sub> compared to CI<sub>1</sub> (Litovsky *et al.*, 2006; Van-Deun *et al.*, 2010; Chadha  
73 *et al.*, 2011). The durations of bilateral implant use in these studies vary from three months to  
74 five years, however no single study has reported changes in SRM over time for the same  
75 children.

76

77 Given the potential influence of auditory system plasticity, it is not straight-forward to predict  
78 the development trajectory of speech discrimination and SRM of sequentially-implanted  
79 children based on data obtained during the first two years post-CI<sub>2</sub>. Knowledge of longer  
80 term outcomes would inform clinicians' management decisions for children with an existing  
81 single cochlear implant, as well as provide realistic expectations for families of such children.  
82 Therefore, this paper presents data from a small scale study conducted at our clinical centre  
83 that describes monaural SRTs in quiet, binaural SRTs in noise and SRM outcomes for  
84 sequentially implanted children at two and four years post-CI<sub>2</sub>.

85

## 86 METHODS

87

88 Data were collected from 17 (eight male, nine female) children who had received sequential  
89 cochlear implants at our clinical service. For inclusion in this study we identified children  
90 who were over four years of age, developmentally able to participate and consistent users of  
91 both CI<sub>1</sub> and CI<sub>2</sub>. We included only children with monaural aided thresholds of 35 dB HL or  
92 better at 0.25, 0.5, 1, 2, 4 and 6 kHz bilaterally. Data were collected for each child at two and  
93 four years post-Cl<sub>2</sub> as part of their routine clinical management. Details regarding each  
94 participating child are given in Table 1. The age range of children at two years post-Cl<sub>2</sub> was  
95 62 to 156 months (median = 119 months) and at 4 years post-Cl<sub>2</sub> was 85 to 182 months  
96 (median = 142 months). The time between Cl<sub>1</sub> and Cl<sub>2</sub> ranged from 19 to 95 months (median  
97 = 49 months). Based on information available in their medical records including audiological  
98 test results, correspondence and parental reports children were assumed to have congenital  
99 profound sensori-neural hearing loss. A number of children were notably older than others at  
100 Cl<sub>1</sub> (i.e. ID 16, 17, 18, 19, 22 and 24) due to a range of non-audiological factors (e.g. repeated  
101 non-attendance at consultations, professional concern regarding family support). Table 1 also  
102 shows the internal implants, external speech processors and processing strategies used by  
103 each child in each ear at both test intervals. For the majority of participants these remained  
104 constant across the time interval. However, two participants (ID 5 and 8) with devices by  
105 Cochlear (Sydney, New South Wales, Australia) had changed from using Freedom<sup>TM</sup> to  
106 CP810<sup>TM</sup> speech processors between assessments and one other participant (ID19) with  
107 devices by MED-EL (Innsbruck, Austria) had changed speech processing strategy from  
108 HDCIS<sup>TM</sup> to FSP<sup>TM</sup> in one ear. Changes in speech processor hardware and processing  
109 strategy can influence speech discrimination (e.g. Kleine Punte *et al.*, 2014, Mosnier *et al.*,  
110 2014.). However, the changes for these three children are considered to be relatively minor

111 and as such will account for only small changes in speech discrimination performance. The  
112 effects of the other characteristics noted in Table 1 are effectively controlled for by the  
113 longitudinal design of this study.

114

115 Measurement of SRT in quiet and noise was achieved using the IHR Automated McCormick  
116 Toy Discrimination Test (Summerfield *et al.*, 1994) presented via the York Crescent of  
117 Sound (Kitterick *et al.*, 2011). The York Crescent of Sound consists of nine Canton Plus  
118 XS.2 loudspeakers (Niederlauken, Germany), each at a height of 1.1 metre, arranged in a  
119 horizontal semi-circle of radius 1.45 metres from +90° (90 ° to the right of the child) to -90°  
120 azimuth (90° to the left of the child). Presentation of speech and noise signals was controlled  
121 via system software and routed to the loudspeakers via a MOTU UltraLite Mk3 (Cambridge,  
122 USA) audio interface and Alesis RA-150 dual-channel amplifiers (Cumberland, USA).

123

124 Speech signals were recorded by Summerfield *et al.* (1994) using a female voice. They  
125 consisted of the introductory phrase “Point to the” followed by the name of one of 10 to 14  
126 toys (phonemically paired e.g. “key” and “tree”) selected at random by system software. The  
127 introductory phrase component of the speech signal had duration of 500 ms. The noise signal  
128 was a burst of broadband (pink) noise with duration of 1400 ms (linear rise-fall = 200 ms;  
129 steady-state = 1000 ms). The noise signal was presented 300 ms following the onset of the  
130 speech signal so that it was at steady-state for the duration of the toy name component of the  
131 speech signal.

132

133 All testing took place in a sound-attenuated room with the child seated so that their head was  
134 an equal distance from all loudspeakers. Children were asked to select which toy name they

135 **Table 1 Participants' characteristics**

Identification code	First CI side	Aetiology	Age confirmed profoundly deaf *\$	Age at first CI *	Age at second CI *	First CI model	Second CI model	Processors at 2 year assessment	Processors at 4 year assessment	1 <sup>st</sup> CI strategy at 2 year assessment	1 <sup>st</sup> CI strategy at 4 year assessment	2 <sup>nd</sup> CI strategy at 2 year assessment	2 <sup>nd</sup> CI strategy at 4 year assessment
5	R	Unknown	13	22	38	CI24 RE(CA)	CI24 R(CA)	Freedom	CP810	ACE, ADRO	ACE, ADRO	ACE, ADRO	ACE, ADRO
6	L	Unknown	0	29	55	Sonata ti100	Sonata ti100	Opus2	Opus2	FSP	FSP	FSP	FSP
8	R	Unknown	11	23	79	CI24R(CA)	CI24 RE(CA)	Freedom	CP810	ACE, ADRO	ACE, ADRO	ACE, ADRO	ACE, ADRO
10	L	Unknown	16	33	59	Pulsar ci 100	Sonata ti 100	Opus2	Opus2	FSP	FSP	FSP	FSP
11	R	Unknown	0	28	78	CI24RE(CA)	CI24RE(CA)	CP810	CP810	ACE	ACE	ACE	ACE
12	R	Unknown	0	17	63	CI24RE(CA)	CI24RE(CA)	CP810	CP810	ACE with ADRO	ACE, ADRO	ACE, ADRO	ACE, ADRO
16	R	Unknown	0	38	59	CI24RE(CA)	CI24RE(CA)	CP810	CP810	ACE, ADRO & auto-sensitivity			
17	R	CMV	48	62	102	CI24RE(CA)	CI24RE(CA)	CP810	CP810	ACE, ADRO	ACE, ADRO	ACE, ADRO	ACE, ADRO
18	R	CMV	51	62	102	CI24RE(CA)	CI24RE(CA)	CP810	CP810	ACE, ADRO	ACE, ADRO	ACE, ADRO	ACE, ADRO
25	R	Unknown	17	22	118	C40+	Sonata ti 100	Opus2	Opus2	FSP	FSP	FSP	FSP
27	R	Usher's syndrome	0	34	129	C40+	Sonata ti 100	Opus2	Opus2	FSP	FSP	FSP	FSP
19	R	Unknown genetic	0	39	105	Pulsar ci 100	Sonata ti 100	Opus2	Opus2	HDCIS	FSP	FSP	FSP
26	L	Usher's syndrome	0	32	93	Pulsar ci 100	Sonata ti 100	Opus2	Opus2	FSP	FSP	FSP	FSP
22	L	Unknown	19	48	98	Pulsar ci 100	Sonata ti 100	Opus2	Opus2	FSP	FSP	FSP	FSP
31	L	Unknown	0	18	37	CI24RE Straight	CI24RE Straight	CP810	CP810	ACE, ADRO	ACE, ADRO	ACE, ADRO	ACE, ADRO
21	R	Unknown	0	33	114	C40+	Sonata ti 100	Opus2	Opus2	FSP	FSP	FSP	FSP
24	R	Unknown genetic	28	58	130	C40+	Sonata ti 100	Opus2	Opus2	FSP	FSP	FSP	FSP

136 \*Ages given in months. \$Where profound loss confirmed on immediate follow-up after failing neonatal hearing screen, age of diagnosis given as 137 0 months. Profound deafness defined as an unaided loss of 90 dB HL or worse at 2 kHz and 4 kHz bilaterally.

138 heard by pointing to a toy on a table in front of them, or selecting an image of the toy on a  
139 touch-screen.

140

141 Monaural SRTs in quiet were assessed first. Speech signals were presented from 0° azimuth  
142 at an initial level of 45 – 55 dB SPL whilst only one cochlear implant was activated. To  
143 encourage compliance with testing, the children were allowed to choose which speech  
144 processor to remove first. A one-down, one-up adaptive procedure with step sizes of 6 dB  
145 was used for the first two reversals, followed by six reversals using a two-down, one-up  
146 adaptive procedure with step sizes of 3 dB. The last six reversals were used to estimate SRT.  
147 The task was then repeated to measure SRT with only the other cochlear implant activated.

148

149 Binaural SRTs in noise were assessed next. First the speech signal and noise were presented  
150 from 0° azimuth ( $S_0N_0$ ) to ensure that one standard outcome of listening in noise was  
151 obtained for each child should they withdraw co-operation before the end of the test session.  
152 Subsequently the speech signal remained at 0° azimuth and the noise was presented from  
153 –90° or +90° azimuth. Both –90° and +90° azimuth result in noise being closest to either  $CI_1$   
154 or  $CI_2$ . This is indicated within this paper by referring to these noise conditions as  $S_0N_{CI1}$  and  
155  $S_0N_{CI2}$  respectively. The speech signal was fixed at 60 dB(A) SPL and the noise signal  
156 varied from an initial level of 30 to 38 dB SPL using an adaptive procedure. The first two  
157 reversals followed a one-down one-up procedure with step sizes of 6 dB. Six further  
158 reversals using a two-down one-up procedure with step sizes of 3 dB were used to establish  
159 SRT in noise, expressed as a signal to noise ratio (SNR). If the noise reached a maximum  
160 level of 60 dB SPL, i.e. a SNR of 0 dB, the speech signal was presented at adaptively quieter  
161 levels in order to adjust the SNR.

162

163 SRM was calculated for each participant by subtracting their SRT in noise for  $S_0N_{CI1}$  and  
164  $S_0N_{CI2}$  from their SRT for  $S_0N_0$ . This resulted in two SRM measurements for each  
165 participant, i.e. SRM with noise located at  $CI_1$  ( $SRM_{CI1}$ ) and noise located at  $CI_2$  ( $SRM_{CI2}$ ).

166

167 Statistical analysis was performed using two-level regression modelling (e.g. Goldstein,  
168 2011; Snijders and Bosker, 2011) with the levels of the model being measurement (within-  
169 participant) and participant (between-participant). For each dependent variable (SRT in  
170 quiet, SRT in noise and SRM) a series of models were used to explore the effect of  
171 explanatory variables (i.e. time post- $CI_2$ , implanted ear and noise location). An advantage of  
172 these models is their ability to incorporate the clustering of data inherent in repeated  
173 measures experimental designs, and avoid violating the assumption of independence of data  
174 that underpins single-level regression methods. Models were estimated by the maximum  
175 likelihood method via an iterative generalised least squares procedure (e.g. Goldstein, 1986).  
176 This allowed an estimate of model deviance to be made. The difference between the  
177 deviance of two models (that differ simply by the addition of explanatory variables) can be  
178 used as a test statistic to determine the effect of the explanatory variables on the dependent  
179 variable. This deviance statistic has a  $\chi^2$  distribution with degrees of freedom equal to the  
180 difference in number of variables included in the two models. In addition, regression  
181 coefficients can be tested for significance via the Wald test (see Snijders and Bosker, 2011).

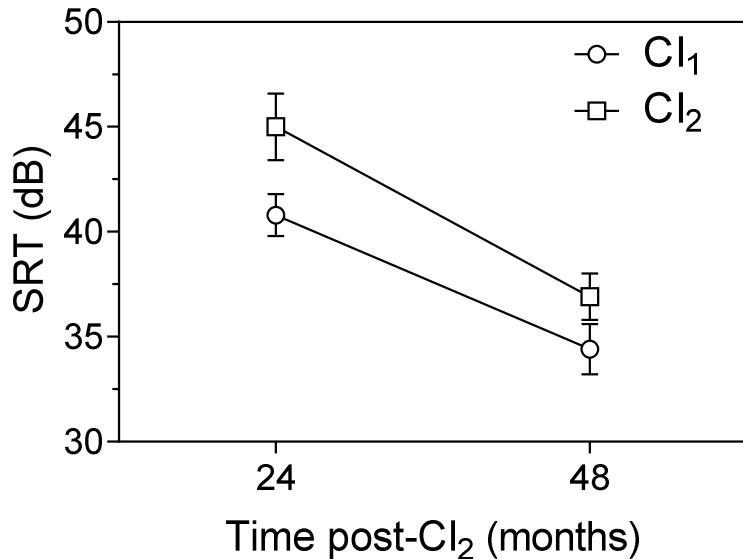
## 182 RESULTS

183

184 Figure 1 shows the mean ( $n = 17$ ) monaural SRTs measured in quiet for CI<sub>1</sub> and CI<sub>2</sub> ears  
185 (circles and squares respectively) at two and four years post-Cl<sub>2</sub>. A number of trends are  
186 clearly evident within the figure. CI<sub>1</sub> ears had lower mean SRT (i.e. better performance) than  
187 CI<sub>2</sub> ears at two years post-Cl<sub>2</sub>. In addition, SRT for both ears reduced (i.e. improved) as a  
188 function of time post-Cl<sub>2</sub>. These observations were confirmed by two-level regression  
189 modelling. Both the inclusion of ear ( $\chi^2 = 5.46, df = 1, p < 0.05$ ) and time post-Cl<sub>2</sub> ( $\chi^2 =$   
190  $37.84, df = 1, p < 0.0001$ ) caused significant reductions in model deviance. Inspection of the  
191 figure also suggests that the improvement in SRT over time was dependent on ear, with a  
192 greater change seen for CI<sub>2</sub> ears (8.1 dB) compared to the CI<sub>1</sub> ears (6.4 dB). However, after  
193 four years post-Cl<sub>2</sub>, CI<sub>1</sub> ears still had lower mean SRT than CI<sub>2</sub> ears. Statistical modelling  
194 including the interaction between ear and time post-second implant showed the difference in  
195 SRT improvement over time to be non-significant ( $\chi^2 = 0.76, df = 1, p = 0.39$ ).<sup>1</sup>

---

<sup>1</sup> For this and all subsequent models reported here, greatest variation was seen at the measurement (within-participant) level, with only minimal variation seen at the participant (between-participant) level. This is in keeping with the longitudinal design of this study. For all models the residuals were confirmed as being normally distributed with mean of zero.



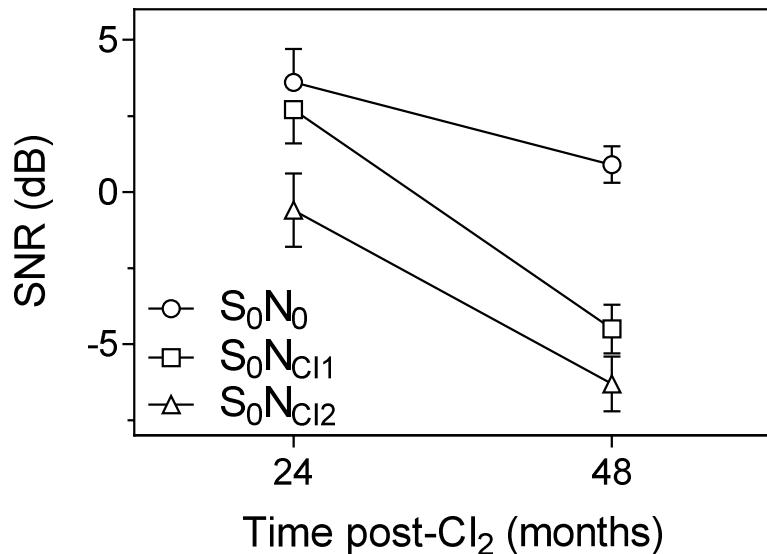
196

197 **Figure 1** Mean monaural SRT in quiet for CI1 (circles) and CI2 (squares) ears as a  
 198 function of time post-Cl2. Error bars represent  $\pm 1$  standard error of the mean (SEM).

199

200 One participant (ID5) had incomplete SRT in noise data and was therefore not included in  
 201 subsequent analysis. The mean ( $n = 16$ ) binaural SRTs measured in noise (expressed as SNR  
 202 in dB) at two and four years post-Cl2 are shown in Figure 2. The figure shows the SNRs  
 203 obtained for the three locations of noise:  $S_0N_0$  (circles),  $S_0N_{CI1}$  (squares) and  $S_0N_{CI2}$   
 204 (triangles). At two and four years post-Cl2, lowest mean SNRs (i.e. better performance) were  
 205 measured at  $S_0N_{CI2}$  with highest SNRs measured at  $S_0N_0$ . For all three noise locations SNRs  
 206 reduced (i.e. improved) as a function of time post-Cl2. The largest improvement was seen at  
 207  $S_0N_{CI1}$  (7.2 dB) followed by  $S_0N_{CI2}$  (5.7 dB), with a smaller improvement (2.7 dB) seen at  
 208  $S_0N_0$ . As a result, mean SRT in noise at  $S_0N_{CI1}$  was most similar to that obtained at  $S_0N_0$  at  
 209 two years but was closest to  $S_0N_{CI2}$  at four years. These observations are confirmed by the  
 210 results of statistical modelling. Both noise location ( $\chi^2 = 25.91$ ,  $df = 2$ ,  $p < 0.0001$ ) and time  
 211 post-Cl2 ( $\chi^2 = 51.30$ ,  $df = 1$ ,  $p < 0.0001$ ) caused highly significant reductions in model  
 212 deviance. The interaction between noise location and time post-Cl2 was also shown to be

213 significant ( $\chi^2 = 10.05$ ,  $df = 2$ ,  $p < 0.01$ ) confirming the difference in improvements seen  
 214 across the three conditions. The model also confirms the convergence of SRT in noise for  
 215  $S_0N_{CI1}$  and  $S_0N_{CI2}$  as a result of the greater improvement seen for  $S_0N_{CI1}$ . Whilst SRT at  
 216  $S_0N_{CI1}$  and  $S_0N_{CI2}$  were significantly different at two years post- $CI_2$  ( $t = 3.27$ ,  $p < 0.001$ ), the  
 217 difference was not significant at four years post- $CI_2$  ( $t = 1.81$ ,  $p = 0.04$ ).<sup>2</sup>

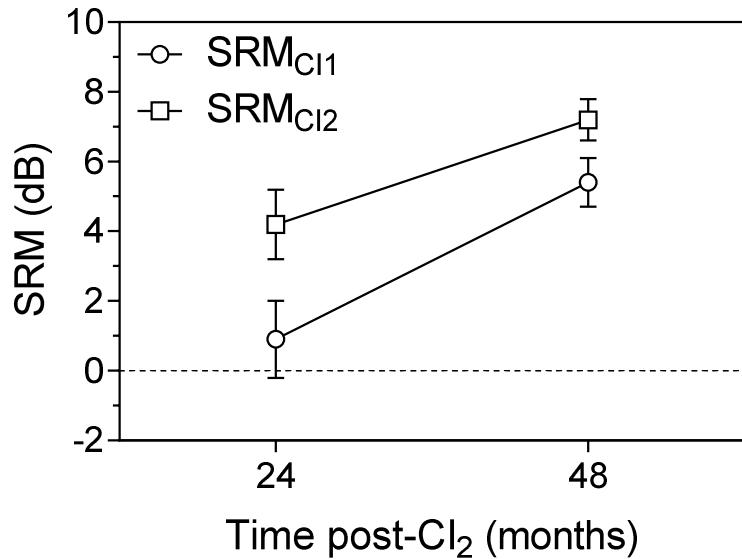


218  
 219 **Figure 2 Mean binaural SRT in noise measured for S0N0 (circles), S0NCI1 (squares)**  
 220 **and S0NCI2 (triangles) as a function of time post-Cl2. Error bars represent  $\pm 1$  SEM.**

221  
 222 Finally, Figure 3 shows the mean ( $n = 16$ ) SRM values obtained as a function of time post-  
 223  $Cl_2$ . SRM values are shown for both noise locations, i.e.  $SRM_{CI1}$  and  $SRM_{CI2}$ . A clear trend  
 224 for both  $SRM_{CI1}$  and  $SRM_{CI2}$  to increase (improve) as a function of time post- $Cl_2$  is evident.  
 225 In addition, a notable difference exists between  $SRM_{CI1}$  and  $SRM_{CI2}$ , with  $SRM_{CI2}$  having  
 226 larger values (i.e. more advantage) than  $SRM_{CI1}$  at two and four years. However, this  
 227 difference becomes smaller as a function of time post- $Cl_2$  from 3.3 dB at two years to 1.8 dB  
 228 at four years. That is,  $SRM_{CI1}$  shows a greater improvement than  $SRM_{CI2}$ , and as a result,

<sup>2</sup> For multiple hypotheses testing a Bonferroni-corrected significance level of  $p < 0.01$  was used.

229 SRM across ears is observed to become more symmetrical over time. Statistical modelling  
 230 confirmed both noise location ( $\chi^2 = 6.34, df = 1, p < 0.05$ ) and time post- $\text{CI}_2$  ( $\chi^2 = 17.00, df =$   
 231 1,  $p < 0.0001$ ) had a significant effect on SRM. The interaction between noise location and  
 232 time was not significant ( $\chi^2 = 0.73, df = 1, p = 0.39$ ), indicating that the time-dependent  
 233 improvements in  $\text{SRM}_{\text{CI}1}$  and  $\text{SRM}_{\text{CI}2}$  were not significantly different.



234

235 **Figure 3** Mean  $\text{SRM}_{\text{CI}1}$  (circles) and  $\text{SRM}_{\text{CI}2}$  (squares) as a function of time post- $\text{Cl}_2$ .  
 236 Error bars represent  $\pm 1$  SEM.

237

238 DISCUSSION

239

240 To date, no longitudinal data have been reported that describe changes in SRM over time for  
241 sequentially-implanted children. Previous investigators (Peters *et al.*, 2007, Sparreboom *et*  
242 *al.*, 2011 and Strom-Roum *et al.*, 2012) have described longitudinal changes in speech  
243 discrimination abilities for this group of children, but these are limited to the first two years  
244 post- $\text{CI}_2$ . The small scale longitudinal study described in this paper is the first to provide a  
245 description of changes in speech discrimination in quiet and noise as well as SRM for  
246 sequentially-implanted children at four years post- $\text{CI}_2$ .

247

248 Our findings demonstrate that the trajectory of improvement in speech discrimination  
249 performance previously reported for up to two years post- $\text{CI}_2$  (Peters *et al.*, 2007;  
250 Sparreboom *et al.*, 2011; Strom-Roum *et al.*, 2012) continues during the next two years. That  
251 is, SRT in both quiet and noise continue to improve for both  $\text{CI}_1$  and  $\text{CI}_2$ . Whilst better  
252 performance is seen for  $\text{CI}_1$ ,  $\text{CI}_2$  shows the greatest improvement over time. This results in  
253 more symmetrical performance across ears.

254

255 Similar findings were also obtained for SRM. Whilst our mean values measured at two years  
256 post- $\text{CI}_2$  were similar to those reported at the same time point by Litovsky *et al.* (2006) and  
257 Sparreboom *et al.* (2011), substantial improvements in SRM for noise presented 90° towards  
258  $\text{CI}_1$  and  $\text{CI}_2$  were observed at four years post- $\text{CI}_2$ . The present data also shows that the  
259 notable asymmetry in SRM evident at two years post- $\text{CI}_2$  (Litovsky *et al.*, 2006; Van-Deun *et*  
260 *al.*, 2010; Chadha *et al.*, 2011) becomes less marked by four years post- $\text{CI}_2$ . However, this  
261 group of sequentially-implanted children did not gain the same symmetrical SRM reported  
262 for simultaneously implanted children at two years post- $\text{CI}_2$  (Chadha *et al.*, 2011).

263

264 In summary, the present findings show that sequentially-implanted children who are  
265 consistent users of two cochlear implants that provide access to sounds at 35 dB HL or better  
266 bilaterally continue to experience substantial improvements in discriminating speech in noise  
267 up to four years post- $CI_2$ , despite the extended period of auditory deprivation in their second-  
268 implanted ear. These findings, along with other evidence (e.g. Smulders *et al.*, 2011) support  
269 the recommendation that children with an existing single implant should be considered for  
270 assessment for a second implant. As a tentative indication of the window of opportunity for  
271 providing a second implant, children in this study who had used a single cochlear implant for  
272 up to 95 months before receiving a second implant still experienced significant improvement  
273 in speech discrimination abilities.

274

275 The increased knowledge of the development of speech discrimination provided by this paper  
276 is useful when counselling families of children considering sequential implantation. As part  
277 of managing expectations families can be made aware of the long time-scale over which  
278 benefits may be obtained. Similarly, some children who have already received a second,  
279 sequential implant struggle to establish consistent use of both devices (Galvin and Hughes,  
280 2012; Fitzgerald *et al.*, 2013). For these families the knowledge that these improvements can  
281 continue beyond two years post- $CI_2$  may serve as motivation to persevere with using the  
282 second cochlear implant and the associated rehabilitation.

283

284 Finally, in order to determine the trajectory of any further changes in speech discrimination  
285 beyond four years post- $CI_2$ , it is recommended that further studies are undertaken with the  
286 aim of measuring speech discrimination performance at longer intervals post- $CI_2$ .

287

288 REFERENCES

289

290 Akeroyd M.A. 2006. The psychoacoustics of binaural hearing. *International Journal of*  
291 *Audiology*, 45(Supplement 1): S25-S33.

292

293 Chadha N.K., Papsin B.C., Jiwani S., Gordon K.A. 2011. Speech detection in noise and  
294 spatial unmasking in children with simultaneous versus sequential bilateral cochlear implants.  
295 *Otology & Neurology*, 32: 1057-1064.

296

297 Fitzgerald M.B., Green J.E., Fang Y., Waltzman S.B. 2013. Factors influencing consistent  
298 device use in pediatric recipients of bilateral cochlear implants. *Cochlear Implants*  
299 *International*, 14(5): 257-265.

300

301 Galvin K.L., Hughes K.C. 2012. Adapting to bilateral cochlear implants: Early post-operative  
302 device use by children receiving sequential or simultaneous implants at or before 3.5 years.  
303 *Cochlear Implants International*, 13(2): 105-112.

304

305 Goldstein, H. 1986. Multilevel mixed linear-model analysis using iterative generalized least-  
306 squares. *Biometrika* 73, 43-56.

307

308 Goldstein, H. 2011. Multilevel statistical models. Chichester: Wiley.

309

310 Gordon K.A., Wong D.D.E., Papsin B.C. 2013. Bilateral input protects the cortex from  
311 unilaterally-driven reorganization in children who are deaf. *Brain*, 136; 1609-1625.

312

313 Green K.M.J., Julyan P.J., Hastings D.L., Ramsden R.T. 2011. Cortical activations in  
314 sequential bilateral cochlear implant users. *Cochlear Implants International*, 12(1): 3-9.

315

316 Hawley M.L., Litovsky R.Y., Culling J.F. 2004. The benefit of binaural hearing in a cocktail  
317 party: Effect of location and type of interferer. *Journal of the Acoustical Society of America*,  
318 115(2): 833-843.

319

320 Kitterick P.T., Lovett R.E.S., Goman A.M., Summerfield A.Q. 2011. The AB-York crescent  
321 of sound: An apparatus for assessing spatial-listening skills in children and adults. *Cochlear  
322 Implants International*, 12(3): 164-169.

323

324 Kleine Punte A., De Bodt M., Van de Heyning P. 2014. Long-Term improvement of speech  
325 perception with the fine structure processing coding strategy in cochlear implants.  
326 *Otorhinolaryngology*, 76: 36-43.

327

328 Litovsky R.Y., Johnstone P.M., Godar S.P. 2006. Benefits of bilateral cochlear implants  
329 and/or hearing aids in children. *International Journal of Audiology*; 45(Suppl 1), S78-S91.

330

331 Mosnier I., Marx M., Venail F., Loudon N., Roux-Vaillard S., Sterkers O. 2014. Benefits  
332 from upgrade to the CP810<sup>TM</sup> sound processor for Nucleus<sup>®</sup> 24 cochlear implant recipients.  
333 *European Archives of Otorhinolaryngology*, 271: 49-57.

334

335 Peters B.R., Litovsky R., Parkinson A., Lake J. 2007. Importance of age and postimplantation  
336 experience on speech perception measures in children with sequential bilateral cochlear  
337 implants. *Otology & Neurotology*, 28: 649-657.

338

339 Sharma A., Gilley P.M., Martin K., Roland P., Bauer P., Dorman M. 2007. Simultaneous  
340 versus sequential bilateral implantation in young children: Effects on central auditory system  
341 development and plasticity. *Audiological Medicine*, 5: 218-223.

342

343 Smulders Y.E., MD; Rinia A.B., Maroeska M.D., Rovers M., van Zanten G.A., Grolman W.  
344 2011. What is the effect of time between sequential cochlear implantations on hearing in  
345 adults and children? A systematic review of the literature. *The Laryngoscope*, 121:1942–  
346 1949.

347

348 Snijders, T.A.A., Bosker, R.J. 2011. Multilevel analysis: An introduction to basic and  
349 advanced multilevel modeling. London: Sage Publications Limited.

350

351 Sparreboom M., Snik A.F.M., Mylanus E.A.M. 2011. Sequential bilateral cochlear  
352 implantation in children: Development of the primary auditory abilities of bilateral  
353 stimulation. *Audiology & Neurotology*, 16: 203-213.

354

355 Sparreboom M., Beynon A.J., Snik A.F.M., Mylanus E.A.M. 2013. Auditory cortical  
356 maturation in children with sequential bilateral cochlear implants. *Otology & Neurotology*,  
357 35: 35-42.

358

359 Strom-Roum H., Laurent C., Wie O.B. 2012. Comparison of bilateral and unilateral cochlear  
360 implants in children with sequential surgery. *International Journal of Pediatric  
361 Otorhinolaryngology*, 76: 95-99.

362

363 Summerfield Q., Palmer A., Foster J., Marshall D., Twomey T. 1994. Clinical evaluation and  
364 test-retest reliability of the IHR-McCormick automated toy discrimination test. *British*  
365 *Journal of Audiology*, 28(3): 165-179.

366

367 Van-Deun L., van Wieringen A., Wouters J. 2010. Spatial speech perception benefits in  
368 young children with normal hearing and cochlear implants. *Ear & Hearing*, 31: 702-713.

369