UNITED STATES
SECURITIES AND EXCHANGE COMMISSION
Washington, D.C. 20549

FORM 6-K

REPORT OF FOREIGN PRIVATE ISSUER PURSUANT TO RULE 13a-16 OR 15d-16
UNDER THE SECURITIES EXCHANGE ACT OF 1934

For the month of September, 2014

Commission File Number: 001-36582

Auris Medical Holding AG

(Exact name of registrant as specified in its charter)

Bahnhofstrasse 21
6300 Zug, Switzerland
(Address of principal executive office)

Indicate by check mark whether the registrant files or will file annual reports under cover of Form 20-F or Form 40-F:

Form 20-F X Form 40-F _____

Indicate by check mark if the registrant is submitting the Form 6-K in paper as permitted by Regulation S-T Rule 101(b)(1):

Yes _____ No X

Indicate by check mark if the registrant is submitting the Form 6-K in paper as permitted by Regulation S-T Rule 101(b)(7):

Yes _____ No X

SIGNATURE

Pursuant to the requirements of the Securities Exchange Act of 1934, the registrant has duly caused this report to be signed on its behalf by the undersigned, thereunto duly authorized.

Auris Medical Holding AG

By: /s/ Thomas Meyer
Name: Thomas Meyer
Title: Chief Executive Officer

Date: September 15, 2014

EXHIBIT INDEX

Exhibit Number	Description
99.1	Unaudited Condensed Consolidated Interim Financial Information
99.2	Management's Discussion and Analysis of Financial Condition and Results of Operations
99.3	Press Release dated September 15, 2014
99.4	Second Quarter 2014 Earnings and Business Update Presentation dated September 15, 2014

Exhibit 99.1

Unaudited Condensed Consolidated Interim Financial Information as of June 30, 2014 and For the Six Months Ended June 30, 2014 and 2013

AURIS MEDICAL HOLDING AG

Condensed Consolidated Interim Statement of Profit or Loss and Other Comprehensive Income (unaudited)
For the Six Months Ended June 30, 2014 and 2013 (in CHF)

	Note	SIX MONTHS ENDED JUNE 30 2014	SIX MONTHS ENDED JUNE 30 2013
Research and development		-8,350,008	-6,384,347
General and administrative		-2,554,288	-732,210
Operating loss		**-10,904,296**	**-7,116,557**
Finance expense		-68,031	-11,366
Finance income		121,144	51,119
Loss before tax		**-10,851,183**	**-7,076,804**
Net loss for the period attributable to owners of the Company		**-10,851,183**	**-7,076,804**
Other comprehensive income:			
Items that will never be reclassified to profit or loss			
Remeasurements of defined benefit liability		-423,811	91,981
Items that are or may be reclassified to profit or loss			
Foreign currency translation differences		2,817	-25,915
Other comprehensive income for the period, net of tax		**-420,994**	**66,066**
Total comprehensive loss for the period attributable to owners of the Company		**-11,272,177**	**-7,010,738**
Basic and diluted loss per share	6	-0.59	-0.52

AURIS MEDICAL HOLDING AG

Condensed Consolidated Interim Statement of Financial Position
As of June 30, 2014 (unaudited), and December 31, 2013
(in CHF)

	Note	June 30, 2014	December 31, 2013
ASSETS			
Non-current assets			
Property and equipment		251,095	195,915
Intangible assets		1,482,520	1,482,520
Total non-current assets		**1,733,615**	**1,678,435**
Current assets			
Other receivables		694,396	524,786
Prepayments		713,889	183,137
Cash and cash equivalents		12,130,954	23,865,842
Total current assets		**13,539,239**	**24,573,765**
Total assets		**15,272,854**	**26,252,200**
EQUITY AND LIABILITIES			
Equity			
Share capital		7,536,510	6,487,130
Share premium		48,232,566	35,608,210
Foreign currency translation reserve		56,812	53,995
Accumulated deficit		-44,295,487	-33,115,689
Total shareholders' equity attributable to owners of the Company		**11,530,401**	**9,033,646**
Non-current liabilities			
Employee benefits	5	745,440	328,342
Deferred tax liabilities		327,637	327,637
Total non-current liabilities		**1,073,077**	**655,979**
Current liabilities			
Convertible loans	4	-	13,711,200
Trade and other payables		1,105,180	954,257
Accrued expenses		1,564,196	1,897,118
Total current liabilities		**2,669,376**	**16,562,575**
Total liabilities		**3,742,453**	**17,218,554**
Total equity and liabilities		**15,272,854**	**26,252,200**

AURIS MEDICAL HOLDING AG

Condensed Consolidated Interim Statement of Cash Flows (unaudited)
For the Six Months Ended June 30, 2014 and 2013 (in CHF)

	Note	SIX MONTHS ENDED JUNE 30 2014	SIX MONTHS ENDED JUNE 30 2013
Cash flows from operating activities			
Net loss		-10,851,183	-7,076,804
Adjustments for:			
Depreciation		34,006	13,562
Unrealized exchange differences		2,835	-25,686
Net interest income		19,750	-2,181
Share based payments	5	95,197	51,325
Employee benefits		-6,713	17,995
Changes in:			
Other receivables		-169,492	-17,091
Prepayments		-530,752	72,325
Trade and other payables		150,901	139,438
Accrued expenses		-332,923	1,110,023
Cash used in operating activities		**-11,588,374**	**-5,717,094**
Cash flows from investing activities			
Purchase of property and equipment		-89,185	-30,233
Sale of financial assets		-	-
Interest received		29,885	12,096
Net cash used in investing activities		**-59,300**	**-18,137**
Cash flows from financing activities			
Proceeds from share capital increase		49,600	24,480,060
Share issuance costs		-136,699	-242,115
Interest paid		-	-9,915
Net cash from financing activities		**-87,099**	**24,228,030**
Net increase/(decrease) in cash and cash equivalents		**-11,734,773**	**18,492,799**
Cash and cash equivalents at beginning of the period		23,865,842	63,967
Net effect of currency translation on cash		-115	614
Cash and cash equivalents at end of the period		**12,130,954**	**18,557,380**

1. Reporting entity

Auris Medical Holding AG (the "Company") is domiciled in Switzerland. The Company's registered address is at Bahnhofstrasse 21, 6300 Zug. These consolidated interim financial information comprise the Company and its subsidiaries (together referred to as the "Group" and individually as "Group entities"). The Company is the ultimate parent of the following Group entities:

- Auris Medical AG, Basel, Switzerland (100%)
- Otolanum AG, Zug, Switzerland (100%)
- Auris Medical Inc., Chicago, United States (100%)
- Auris Medical Ltd., Dublin, Ireland (100%)

On April 22, 2014, we changed our name from Auris Medical AG to Auris Medical Holding AG and transferred our operational business to our newly incorporated subsidiary Auris Medical AG, which is now our main operating subsidiary. The Group is primarily involved in the development of pharmaceutical products for the treatment of inner ear disorders, in particular tinnitus and hearing loss. Its most advanced projects are in the late stage of clinical development.

2. Basis of preparation

This interim financial information has been prepared using the same accounting policies and methods of computation as compared with the most recent annual financial statements. The financial information included here-in is not intended to fully comply with IFRS, and in particular with IAS 34. New accounting standards, which have been applied in 2014 for the first time, did not have an impact on the company's financial position, results or cash flows.

The financial information included in this quarterly press release and shareholder information has not been reviewed or audited by our auditors. Quarter results are not necessarily indicative of results to be expected for the full year.

The Group has not early adopted any standard, interpretation or amendment that was issued but is not yet effective.

3. Capital and reserves

Share capital

The issued share capital of Auris Medical Holding AG consisted of:

	AS OF JUNE 30, 2014		AS OF DECEMBER 31, 2013	
	NUMBER	CHF	NUMBER	CHF
Common shares with a nominal value of CHF 0.40 each	88,100	35,240	72,600	29,040
Preferred shares Series A with a nominal value of CHF 0.40 each	5,999,750	2,399,900	5,999,750	2,399,900
Preferred shares Series B with a nominal value of CHF 0.40 each	5,509,100	2,203,640	5,509,100	2,203,640
Preferred shares Series C with a nominal value of CHF 0.40 each	7,244,325	2,897,730	4,636,375	1,854,550
	18,841,275	**7,536,510**	**16,217,825**	**6,487,130**

All shares are fully paid in. All disclosed numbers and nominal value of shares in these interim financial statements are adjusted for the 25:1 stock split effected in December 2013 unless otherwise indicated.

	Common shares		Preferred shares	
	2014	2013	2014	2013
As of January 1	**72,600**	**72,600**	**16,145,225**	**11,508,850**
Common shares with a nominal value of CHF 0.40 each	15,500	-	2,607,950	-
As at June 30, 2014, and June 30, 2013	**88,100**	**72,600**	**18,753,175**	**11,508,850**

Issue of common shares upon exercise of options

On January 21, 2014, three beneficiaries of Option Plan A exercised their right to acquire common shares of the Company at CHF 3.20 per share. This resulted in an increase in the number of outstanding common shares of 15,500 and an increase in the share capital of CHF 6,200. Total proceeds from the exercise to the company were CHF 49,600.

4. Convertible loans

On December 9, 2013, the Company issued non-interest bearing convertible loans to two shareholders with a nominal value of CHF 13,769,976 and a maximum term of 12 months. On January 27, 2014, the loans were converted into new registered Series C shares with nominal value of CHF 0.40 each for CHF 5.28 per share. The company issued 2,607,950 Series C shares with a nominal value of CHF 0.40 each.

On the conversion date of the loan, the liability was derecognized and CHF 1,043,180 was recognized as share capital and CHF 12,717,655 as share premium.

	JUNE 30, 2014
Convertible loans as at December 31	**13,711,200**
Loss on derecognition	9,141
Imputed interest expense for the period	49,635
Derecognition of liability at conversion into equity	13,769,976
Convertible loans at June 30, 2014	**-**

5. Employee benefits

	SIX MONTHS ENDED JUNE 30, 2014	SIX MONTHS ENDED JUNE 30, 2013
Salaries	980,518	356,110
Pension costs	59,547	39,387
Other social benefits	82,149	30,425
Share option cost	95,197	51,325
Other employee cost	97,871	44,774
Total employee benefits	**1,315,280**	**522,021**

6. Loss per share

	SIX MONTHS ENDED JUNE 30, 2014	SIX MONTHS ENDED JUNE 30, 2013
Loss attributable to owners of the Company	-10,851,183	-7,076,804
Weighted average number of shares outstanding	18,448,274	13,616,303
Basic and diluted loss per share	**-0.59**	**-0.52**

For the periods ended June 30, 2014 and 2013 basic and diluted loss per share is based on the weighted average number of shares issued and outstanding and excludes shares to be issued under the stock option plans (Note 7) and, for the 2013 period only, conversion rights related to the convertible loans (Note 8) as they would be anti-dilutive. In case the Group shows a profit in the future, the options may have a dilutive effect on earnings per share and will need to be included in the above calculation.

7. Related party transactions

In January 2014 the convertible loan lenders exercised their conversion option, and the total loan amount of CHF 13,769,976 was converted into 2,607,950 Series C shares of the Company (See Note 8).

The service agreement with Altamira Pharma GmbH, a company fully owned by the CEO, was terminated as of January 31, 2014 with a final payment of CHF 14,500. From 2011, the CEO had been compensated under this agreement. The Company entered into an employment contract with the CEO effective January 1, 2014.

8. Events after the balance sheet date

On August 6, 2014, the underwriters for the Company's IPO subscribed to 9,400,000 shares at USD 6.00 per share yielding gross proceeds (before underwriting fees and IPO costs) of USD 56.4 million. The shares started trading under the ticker symbol EARS on the NASDAQ Global Market on August 7, 2014.

On August 13, 2014, the underwriters exercised their overallotment option for 713,235 shares of the Company, resulting in gross proceeds of USD 4.3m

The combined net proceeds after underwriting fees are USD 56.4 million. Following the IPO and the issue of shares under the underwriter overallotment option, there are 28,954,510 common shares of the Company outstanding.

MANAGEMENT'S DISCUSSION AND ANALYSIS OF FINANCIAL CONDITION AND RESULTS OF OPERATIONS

This management's discussion and analysis is designed to provide you with a narrative explanation of our financial condition and results of operations. We recommend that you read this in conjunction with our unaudited condensed consolidated interim financial information as of June 30, 2014 and for the six months ended June 30, 2014 and 2013 included as Exhibit 1 to this Report on Form 6-K. We also recommend that you read our management's discussion and analysis and our audited consolidated financial statements and unaudited condensed consolidated interim financial statements, and the notes thereto, which appear in our prospectus (our "Final Prospectus") relating to our Registration Statement on Form F-1, as amended (Registration No. 333-197105), filed with the U.S. Securities and Exchange Commission (the "SEC") pursuant to Rule 424(b) under the U.S. Securities Act of 1933, as amended.

Unless otherwise indicated or the context otherwise requires, all references to "Auris Medical" or the "company," "we," "our," "ours," "us" or similar terms refer to Auris Medical AG and its subsidiaries prior to the completion of our corporate reorganization in connection with our initial public offering, and Auris Medical Holding AG and its subsidiaries as of the completion of our corporate reorganization and thereafter. See "Prospectus Summary—Corporate Information" in the Final Prospectus

We prepare and report our consolidated financial statements and financial information in accordance with International Financial Reporting Standards ("IFRS") as issued by the International Accounting Standards Board (the "IASB"). None of our financial statements were prepared in accordance with generally accepted accounting principles in the United States. We maintain our books and records in Swiss Francs. We have made rounding adjustments to some of the figures included in this management's discussion and analysis. Accordingly, numerical figures shown as totals in some tables may not be an arithmetic aggregation of the figures that precede them. Unless otherwise indicated, all references to currency amounts in this discussions and analysis are in Swiss Francs.

This discussion and analysis is dated as of September 15, 2014.

Overview

We are a clinical-stage biopharmaceutical company focused on the development of novel products for the treatment of inner ear disorders. Our most advanced product candidate, AM-101, is in Phase 3 clinical development for acute inner ear tinnitus under a special protocol assessment, or SPA, from the FDA. In two recently completed Phase 2 clinical trials, AM-101 demonstrated a favorable safety profile and statistically significant improvement in tinnitus loudness and other patient reported outcomes. We expect to have top-line Phase 3 clinical data for AM-101 in early 2016. We are also developing AM-111 for acute inner ear hearing loss (acute sensorineural hearing loss, or ASNHL). Following feedback from a pre-IND meeting with the FDA in September 2014, we plan to finalize the design of the late-stage AM-111 clinical program. Implementation of the late-stage AM-111 clinical program may require additional funding. Both acute inner ear tinnitus and hearing loss are conditions for which there is high unmet medical need, and we believe that we have the potential to be the first to market in these indications.

To date, we have financed our operations through our initial public offering of our common shares, private placements of equity securities and short term loans. We have no products approved for commercialization and have never generated any revenues from royalties or product sales. Based on our current plans, we do not expect to generate royalty or product revenues unless and until we obtain marketing approval for, and commercialize, AM-101, AM-111 or any of our other product candidates.

Since inception, we have incurred significant operating losses. We incurred net losses (defined as net losses attributable to the owners of the company) of CHF 15.0 million and CHF 4.6 million for the years ended December 31, 2013 and 2012, respectively and for the six months ended June 30, 2014 and 2013, we incurred net losses of CHF 10.9 million and CHF 7.1 million, respectively. As of June 30, 2014,

we had an accumulated deficit of CHF 44.30 million. We expect to continue incurring losses as we continue our clinical and preclinical development programs, apply for marketing approval for our product candidates and, subject to obtaining regulatory approval of our product candidates, build a sales and marketing force in preparation for the potential commercialization of our product candidates.

Recent Developments

On August 11, 2014, we completed our initial public offering of common shares pursuant to a Registration Statement on Form F-1, as amended (Registration No. 333-197105) that was declared effective on August 5, 2014. Under the registration statement, we sold an aggregate of 10,113,235 common shares, which included 713,235 common shares sold on August 19, 2014 pursuant to an over-allotment option granted to the underwriters. All of these common shares were sold at a price to the public of $6.00 per share. The offering expenses, not including the underwriting discounts and commissions, are estimated at $2.4 million and are payable by us. The offering expenses include SEC registration fees, FINRA filing fees, Nasdaq listing fees and expenses, legal fees and expenses, printing expenses, transfer agent fees and expenses, accounting fees and expenses as well as other miscellaneous fees and expenses. Our common shares are listed on the NASDAQ Global Market under the symbol "EARS."

Collaboration and License Agreements

There have been no material changes to our collaboration and license agreements from those reported in "Management's Discussion and Analysis of Financial Condition and Results of Operations–Collaboration and License Agreements" in the Final Prospectus.

Research and Development Expense

Our research and development expense is highly dependent on the development phases of our research projects and therefore may fluctuate substantially from period to period. Our research and development expense mainly relates to the following key programs:

- AM-101. We have commenced a Phase 3 program of AM-101 comprising two Phase 3 clinical trials (TACTT2 and TACTT3) as well as two open label extension studies (AMPACT1 and AMPACT2). We expect top-line data from the TACTT trials in early 2016. We anticipate that our research and development expenses will increase substantially in connection with these clinical trials.

- AM-111. Following feedback from a pre-IND meeting with the FDA in September 2014, we plan to finalize the design of our late-stage AM-111 clinical program for ASNHL. Implementation of the late-stage AM-111 clinical program may require additional funding . We anticipate that our research and development expenses will increase substantially with the continuation of AM-111's clinical development program.

- Other development programs. Other research and development expenses mainly relate to our preclinical studies of AM-102 and AM-123. The expenses mainly consist of costs for production of the preclinical compounds and costs paid to academic research institutions in conjunction with preclinical testing.

For a discussion of our other key financial statement line items, please see "Management's Discussion and Analysis of Financial Condition and Results of Operations–Financial Operations Overview" in the Final Prospectus.

Results of Operations

The discussion below should be read along with our unaudited condensed consolidated interim financial information as of June 30, 2014 and for the six months ended June 30, 2013 and 2014, and is qualified in its entirely by reference to them.

Comparison of the three months ended June 30, 2014 and 2013

	Three months ended June 30,		
	2014	2013	Change
	(in thousands of CHF)		%
Research and development	(4,201)	(4,411)	(5)%
General and administrative	(845)	(462)	83%
Operating loss	**(5,046)**	**(4,873)**	**4%**
Finance expense	(17)	(1)	
Finance income	71	27	**164%**
Loss before tax	**(4,991)**	**(4,847)**	**3%**
Net loss attributable to owners of the company	**(4,991)**	**(4,847)**	**3%**
Other comprehensive income:			
Items that will never be reclassified to profit or loss			
Remeasurements of defined benefits liability	(407)	49	
Items that are or may be reclassified to profit or loss			
Foreign currency translation differences	(3)	4	
Other comprehensive income	**(410)**	**53**	
Total comprehensive loss attributable to owners of the company	**(5,401)**	**(4,794)**	**13%**

Research and development expense

Research and development expense	Three months ended June 30,		
	2014	2013	Change
	(in thousands of CHF)		%
Clinical Projects	(2,757)	(3,235)	(15)%
Pre-clinical projects	(587)	(301)	95%
Drug manufacture and substance	(159)	(665)	(76)%
Employee benefits	(428)	(125)	242%
Other research and development expenses	(269)	(85)	217%
Total	**(4,201)**	**(4,411)**	**(5)**%

Research and development expense decreased 5% from CHF 4.4 million in the three months ended June 30, 2013 to CHF 4.2 million in the three months ended June 30, 2014. The variances in expense between the three months ended June 30, 2013 and the corresponding period in 2014 are mainly due to the following projects:

- *Clinical Projects.* In the three months ended June 30, 2014 we incurred lower clinical expenses than in the three months ended June 30, 2013 primarily due to fluctuations in upfront and milestone costs charged by contracted service providers.

- *Pre-clinical projects*. In the three months ended June 30, 2014, pre-clinical expenses were higher than in the three months ended June 30, 2013 as additional studies for AM-111 were ongoing.

- *Drug manufacture and substance.* In the three months ended June 30, 2014 we incurred lower costs related to the manufacture, and storage of supplies for clinical material compared to the three months ended June 30, 2013, primarily due to fluctuations in the timing of raw material purchases and the manufacture of clinical trial supplies.

- *Employee Benefits*. Employee expenses were significantly higher in the three months ended June 30,2014 than in the three months ended June 30, 2013 due to a higher headcount.

General and administrative expense

General and administrative expense increased 83% from CHF 0.5 million in the three months ended June 30, 2013 to CHF 0.9 million in the three months ended June 30, 2014. The increase was primarily related to higher legal and auditing expenses in connection with preparation for the IPO.

We expect that general and administrative expense will increase in the future as our business expands and we incur additional costs associated with operating as a public company.

Finance income

Finance income increased 164% to CHF 0.1 million in the three months ended June 30, 2014. Finance income in these periods consisted primarily of interest income recognized on short-term deposits.

Finance income increased in the three months ended June 30, 2014 as compared to the three months ended June 30, 2013 as a result of higher unrealized exchange gains.

Finance expense

Finance expense was insignificant in the three months ended June 30, 2014 and 2013.

Remeasurements of defined benefits liability

Remeasurements of the net defined benefit liability, which comprise actuarial gains and losses, the return on plan assets (excluding interest) and the effect of the asset ceiling (if any, excluding interest), decreased from CHF 0.1 million in the three months ended June 30, 2013 to CHF (0.4) million in the three months ended June 30, 2014.

Foreign currency translation differences

Foreign currency translation differences were insignificant in the three months ended June 30, 2014 and 2013.

Comparison of the six months ended June 30, 2014 and 2013

	Six months ended June 30,		
	2014	**2013**	**Change**
	(in thousands of CHF)		**%**
Research and development	(8,350)	(6,384)	31%
General and administrative	(2,554)	(732)	248%
Operating loss	**(10,904)**	**(7,117)**	**53%**
Finance expense	(68)	(11)	499%
Finance income	121	51	137%
Loss before tax	**(10,851)**	**(7,077)**	**53%**
Net loss attributable to owners of the company	**(10,851)**	**(7,077)**	**53%**
Other comprehensive income:			
Items that will never be reclassified to profit or loss			
Remeasurements of defined benefits liability	(424)	92	
Items that are or may be reclassified to profit or loss			
Foreign currency translation differences	3	(26)	
Other comprehensive income	**(421)**	**66**	
Total comprehensive loss attributable to owners of the company	**(11,272)**	**(7,011)**	**61%**

Research and development expense

Research and development expense	Six months ended June 30, 2014	2013	Change
	(in thousands of CHF)		%
Clinical Projects	(5,161)	(4,298)	20%
Pre-clinical projects	(1,152)	(701)	64%
Drug manufacture and substance	(784)	(894)	(12)%
Employee benefits	(797)	(349)	129%
Other research and development expenses	(455)	(143)	218%
Total	**(8,350)**	**(6,384)**	**31%**

Research and development expense increased 31% from CHF 6.4 million in the six months ended June 30, 2013 to CHF 8.4 million in the six months ended June 30, 2014. The variances in expense between the six months ended June 30, 2013 and the corresponding period in 2014 are mainly due to the following projects:

- *Clinical Projects.* In the six months ended June 30, 2014 we incurred higher clinical expenses due to the progression of our AM-101 Phase III clinical development program. In addition, we conducted preparatory activities related to our late stage AM-111 clinical program, whereas in the six months ended June 30, 2013 the Phase II had been completed with a concomitant decline in spending.

- *Pre-clinical projects*. In the six months ended June 30, 2014, pre-clinical expenses were higher than in the six months ended June 30, 2013 as more pre-clinical projects were ongoing related to the AM-101 and AM-111 projects.

- *Drug manufacture and substance.* In the six months ended June 30, 2014 we continued to incur substantial costs related to the manufacture, packaging and distribution of clinical trial supplies. However, expense levels decreased compared to the six months ended June 30, 2013 primarily due to fluctuations in the timing of purchases of raw material and the manufacture of clinical trial supplies.

- *Employee Benefits*. Employee expenses were significantly higher in the six months ended June 30, 2014 than in the six months ended June 30, 2013 due to a higher headcount.

General and administrative expense

General and administrative expense increased more than threefold from CHF 0.7 million in the six months ended June 30, 2013 to CHF 2.6 million in the six months ended June 30, 2014. The increase was primarily related to higher legal and auditing expenses in relation to the IPO preparations.

We expect that general and administrative expense will increase in the future as our business expands and we incur additional costs associated with operating as a public company.

Finance income

Finance income increased 137% to CHF 0.1 million in the six months ended June 30, 2014. Finance income in these periods consisted primarily of unrealized exchange gains and interest income recognized on short-term deposits.

Finance income increased in the six months ended June 30, 2014 as compared to the six months ended June 30, 2013 as a result of an increase in average cash and cash equivalents following our private placement of preferred equity securities to new and existing investors and the issuance of a convertible

bond to existing investors with total net proceeds of CHF 24.1 million and CHF 13.8 million, in April and December 2013, respectively.

Finance expense

Finance expense increased fivefold to CHF 0.1 million in the six months ended June 30, 2014. Higher finance expenses were mainly due to an increase in interest expenses due to the issuance to existing shareholders of the convertible bond in a principal amount of CHF 13.8 million in December 2013.

Remeasurements of defined benefits liability

Remeasurements of the net defined benefit liability, which comprise actuarial gains and losses, the return on plan assets (excluding interest) and the effect of the asset ceiling (if any, excluding interest), decreased from CHF 0.1 million in the six months ended June 30, 2013 to (CHF 0.4) million in the six months ended June 30, 2014.

Foreign currency translation differences

Foreign currency translation differences were insignificant in the six months ended June 30, 2014 and 2013.

Liquidity and Capital Resources

Since inception, we have incurred significant operating losses. To date, we have not generated any product sale revenue. We have financed our operations primarily through our initial public offering of our common shares, private placements of equity securities and loans from existing shareholders.

Cash flows

Comparison of the three months ended June 30, 2014 and 2013

The table below summarizes our consolidated statement of cash flows for the three months ended June 30, 2014 and 2013:

	Three months ended June 30,	
	2014	**2013**
	(in thousands of CHF)	
Cash used in operating activities	(6,402)	(4,208)
Net cash used in investing activities	(3)	(7)
Net cash from financing activities	-	22,750
Net effect of currency translation on cash	0	0
Cash and cash equivalents at the beginning of the period	18,535	21
Cash and cash equivalents at the end of the period	**12,131**	**18,557**

The increase in cash used in operating activities by 52% from CHF 4.2 million in the three months ended June 30, 2013 to CHF 6.4 million in the three months ended June 30, 2014 was mainly due to higher research and development and general and administrative expenses.

Net cash used in investing activities was insignificant in the three months ended June 30, 2014 and 2013.

The decrease in net cash from financing activities from CHF 22.7 million in the three months ended June 30, 2013 to CHF nil in the three months ended June 30, 2014 reflects completion of the first tranche of our Series C financing in April 2013.

Comparison of the six months ended June 30, 2014 and 2013

The table below summarizes our consolidated statement of cash flows for the six months ended June 30, 2014 and 2013:

	Six months ended June 30,	
	2014	**2013**
	(in thousands of CHF)	
Cash used in operating activities	(11,588)	(5,717)
Net cash used in investing activities	(59)	(18)
Net cash from financing activities	(87)	24,228
Net effect of currency translation on cash	(0)	1
Cash and cash equivalents at the beginning of the period	23,866	64
Cash and cash equivalents at the end of the period	**12,131**	**18,557**

The increase in cash used in operating activities by 103% from CHF 5.7 million in the six months ended June 30, 2013 to CHF 11.6 million in the six months ended June 30, 2014 was mainly due to higher research and development and general and administrative expenses.

The increase in net cash used in investing activities by 227 % to CHF 0.1 million in the six months ended June 30, 2014 was due to higher investment in manufacturing equipment partially offset by higher interest income due to a higher cash base.

The decrease in net cash from financing activities from CHF 24.2 million in the six months ended June 30, 2013 to CHF (0.1) million in the six months ended June 30, 2014 reflects the completion of the first tranche of the Series C financing in April 2013 as well as stamp duties paid on the conversion of the convertible loan and exercise of stock options in January 2014, which was partially offset by the proceeds received through the exercise of said stock options.

Cash and funding sources

During the three and six months ended June 30, 2014, we did not obtain new financing except from the exercise of stock options. As of June 30, 2014, we had no long term debt and had no ongoing material financial commitments, such as lines of credit or guarantees that are expected to affect our liquidity over the next five years, other than leases.

Funding requirements

We believe that our existing cash and cash equivalents will enable us to fund our operating expenses and capital expenditure requirements for at least 24 months. We have based this estimate on assumptions that may prove to be wrong, and we could use our capital resources sooner than we currently expect. Our future funding requirements will depend on many factors, including but not limited to:

- the scope, rate of progress, results and cost of our clinical trials, nonclinical testing, and other related activities;
- the cost of manufacturing clinical supplies, and establishing commercial supplies, of our product candidates and any products that we may develop;
- the number and characteristics of product candidates that we pursue;
- the cost, timing, and outcomes of regulatory approvals;
- the cost and timing of establishing sales, marketing, and distribution capabilities; and
- the terms and timing of any collaborative, licensing, and other arrangements that we may establish, including any required milestone and royalty payments thereunder.

We expect that we will require additional funding to advance our late-stage AM-111 clinical program. In addition, we expect that we will require additional capital to commercialize our product candidates AM-101 and AM-111. If we receive regulatory approval for AM-101 or AM-111, and if we choose not to grant any licenses to partners, we expect to incur significant commercialization expenses related to product manufacturing, sales, marketing and distribution, depending on where we choose to commercialize. We also expect to incur additional costs associated with operating as a public company following this offering. Additional funds may not be available on a timely basis, on favorable terms, or at all, and such funds, if raised, may not be sufficient to enable us to continue to implement our long-term business strategy. If we are not able to raise capital when needed, we could be forced to delay, reduce or eliminate our product development programs or commercialization efforts.

For more information as to the risks associated with our future funding needs, see "Risk Factors" in the Final Prospectus.

Contractual Obligations and Commitments

There have been no material changes to our contractual obligations outside the ordinary course of our business from those reported in "Management's Discussion and Analysis of Financial Condition and Results of Operations – Contractual Obligations and Commitments" in the Final Prospectus.

Off-balance sheet arrangements

As of the date of this discussion and analysis, we do not have any, and during the periods presented we did not have any, off-balance sheet arrangements except for the Operating Lease mentioned in "Management's Discussion and Analysis of Financial Condition and Results of Operations – Contractual Obligations and Commitments" in the Final Prospectus.

Significant Accounting Policies and Use of Estimates and Judgment

There have been no material changes to the significant accounting policies and estimates described in "Management's Discussion and Analysis of Financial Condition and Results of Operations – Significant accounting policies and use of estimates and judgment" in the Final Prospectus.

Recent Accounting Pronouncements

Except for IFRS 9 for which the impact cannot be determined with sufficient reliability, there are no IFRS standards as issued by the IASB or interpretations issued by the IFRS interpretations committee that

are effective for the first time for the financial year beginning on or after January 1, 2014 that would be expected to have a material impact on our financial position.

JOBS Act Exemption

On April 5, 2012, the JOBS Act was signed into law. The JOBS Act contains provisions that, among other things, reduce certain reporting requirements for an "emerging growth company." As an emerging growth company, we are not required to provide an auditor attestation report on our system of internal controls over financial reporting. This exemption will apply for a period of five years following the completion of our initial public offering or until we no longer meet the requirements of being an "emerging growth company," whichever is earlier. We would cease to be an emerging growth company if we have more than $1.0 billion in annual revenue, have more than $700 million in market value of our common shares held by non-affiliates or issue more than $1.0 billion of non-convertible debt over a three-year period.

Cautionary Statement Regarding Forward Looking Statements

Forward-looking statements appear in a number of places in this discussion and analysis and include, but are not limited to, statements regarding our intent, belief or current expectations. Many of the forward-looking statements contained in this discussion and analysis can be identified by the use of forward-looking words such as "anticipate," "believe," "could," "expect," "should," "plan," "intend," "estimate" and "potential," among others. Forward-looking statements are based on our management's beliefs and assumptions and on information currently available to our management. Such statements are subject to risks and uncertainties, and actual results may differ materially from those expressed or implied in the forward-looking statements due to of various factors, including, but not limited to, those identified under the section entitled "Risk Factors" in the Final Prospectus. These risks and uncertainties include factors relating to:

- our operation as a development stage company with limited operating history and a history of operating losses;
- our need for substantial additional funding before we can expect to become profitable from sales of our products;
- our dependence on the success of AM-101 and AM-111, which are still in clinical development and may eventually prove to be unsuccessful;
- the chance that we may become exposed to costly and damaging liability claims resulting from the testing of our product candidates in the clinic or in the commercial stage;
- uncertainty surrounding whether any of our product candidates will receive regulatory approval, which is necessary before they can be commercialized;
- if our product candidates obtain regulatory approval, our being subject to expensive ongoing obligations and continued regulatory overview;
- enacted and future legislation may increase the difficulty and cost for us to obtain marketing approval and commercialization;
- the chance that we do not obtain orphan drug exclusivity for AM-111, which would allow our competitors to sell products that treat the same conditions;
- dependence on governmental authorities and health insurers establishing adequate reimbursement levels and pricing policies;

- our products may not gain market acceptance, in which case we may not be able to generate product revenues;
- our reliance on our current strategic relationships with INSERM or Xigen and the potential failure to enter into new strategic relationships;
- our reliance on third parties to conduct our nonclinical and clinical trials and on third-party single-source suppliers to supply or produce our product candidates; and
- other risk factors discussed under “Risk Factors” included in the Final Prospectus

Forward-looking statements speak only as of the date they are made, and we do not undertake any obligation to update them in light of new information or future developments or to release publicly any revisions to these statements in order to reflect later events or circumstances or to reflect the occurrence of unanticipated events.

Auris Medical News Release

Auris Medical Holding AG Reports Second Quarter 2014 Financial Results and Provides Business Update

Zug, Switzerland, September 15, 2014 –Auris Medical Holding AG (NASDAQ: EARS) today provided an update on the Company's business and announced financial results for the second quarter ended June 30, 2014.

Recent Business Highlights

- Auris Medical raised net proceeds (after underwriting discounts) of approximately $56.4 million in its initial public offering (IPO), selling 10,113,235 of its common shares, including 713,235 common shares sold pursuant to the underwriters' exercise of their over-allotment option. On August 6, 2014, Auris Medical's shares began trading on the NASDAQ Global Market under the symbol "EARS".

- The AM-101 Phase 3 clinical program in acute inner ear tinnitus remains on track. Ramp-up of clinical trial sites in North America and Europe progressed with approximately 100 sites initiated at the end of August 2014. As previously announced, Auris Medical expects to enroll a total of 930 patients: 630 in the acute stage (i.e. up to three months from onset) and 300 in the post acute stage (i.e. onset between 4 and 12 months). The first patients that completed the TACTT trials rolled over into the open label AMPACT trials in April and May 2014.

- In May 2014, Auris Medical presented data on the determination of clinically meaningful change in subjective tinnitus loudness at the XI. International Tinnitus Seminar in Berlin, Germany. Anchor- and distribution-based statistical analyses of clinical data support the use of a 2 point reduction on a 0-10 numerical rating scale as a responder definition. Improvement in subjective tinnitus loudness is the primary endpoint in the TACTT clinical trials.

- Auris Medical filed additional patent applications in the US relating to the use of inhibitors of the c-Jun N-Terminal Kinase (JNK) such as AM-111 for the treatment of tinnitus or Menière's Disease. In addition the Company filed a continuation to its earlier US application relating to the use of polymer-based formulations in the treatment of disorders of the middle and inner ear.

- Preparations for the AM-111 late stage clinical program in ASHNL are underway. Following discussions with the European Medicines Agency on the design for a pivotal Phase 3 trial, Auris Medical expects further regulatory feedback from a pre-IND meeting with the FDA in September 2014. Based on that feedback, the Company expects to finalize the design of the late-stage AM-111 clinical program.

Auris Medical Holding AG · Bahnhofstrasse 21 · CH-6300 Zug · Tel.+41 41 729 71 94 · www.aurismedical.com

"During the second quarter Auris Medical has made significant progress, both financially and operationally," said Thomas Meyer, Auris Medical's founder, Chairman and CEO. "With the successful completion of our recent IPO, we have obtained the means to complete the Phase 3 clinical development of AM-101 for the treatment of patients with acute inner ear tinnitus and to further advance our other research and development programs."

Financial Results

As of June 30, 2014, the Company had CHF 12.1 million in cash and cash equivalents. Operating expenses for the three months ended June 30, 2014 were CHF 5.0 million, with CHF 4.2 million attributable to research and development. This compares to operating expenses of CHF 4.9 million and research and development expenses of CHF 4.4 million for the same period in 2013. The Company reported a net loss for the quarter ended June 30, 2014 of CHF 5.0 million, or CHF 0.27 per share. This compares to a net loss of CHF 4.8 million, or CHF 0.33 per share, for the same period in 2013.

For the six month period ended June 30, 2014, operating expenses were CHF 10.9 million, with CHF 8.4 million attributable to research and development. This compares to operating expenses of CHF 7.1 million and research and development expenses of CHF 6.4 million for the same period in 2013. The Company reported a net loss for the six months ended June 30, 2014 of CHF 10.9 million, or CHF 0.59 per share. This compares to a net loss of CHF 7.1 million, or CHF 0.52 per share, for the same period in 2013.

The increases in operating expenses, and resulting increases in net loss, for the three- and six-month periods ended June 30, 2014 over the comparable periods in 2013 reflect primarily the progression of the AM-101 Phase 3 clinical development program, preparations for the late stage AM-111 clinical program, headcount expansion and higher legal and auditing expenses related to the IPO preparations.

Conference Call / Webcast Information

Auris Medical will host a live conference call and webcast to discuss the Company's financial results and provide a general business update. The call is scheduled for September 15, 2014 at 8:00 a.m. Eastern Time. To participate in this conference call, dial 1877 280 2342 (USA) or +1 718 354 1158 (International), and enter passcode 3215731. A live, listen-only audio webcast of the conference call can be accessed on the Investor Relations section of the Auris Medical website at: www.aurismedical.com. A replay will be available approximately two hours following the live call also on the Company's website.

About Auris Medical

Auris Medical is a Swiss biopharmaceutical company dedicated to developing therapeutics that address important unmet medical needs in otolaryngology. The Company is currently focusing on the development of treatments for acute inner ear tinnitus (AM-101) and for acute inner ear hearing loss (AM-111) by way of intratympanic injection with biocompatible gel formulations. In addition, Auris Medical is pursuing early-stage research and development projects. The Company was founded in 2003 and is headquartered in Zug, Switzerland.

Forward-looking Statements

This press release may contain statements that constitute "forward-looking statements" within the meaning of Section 27A of the Securities Act of 1933 and Section 21E of the Securities Exchange Act of 1934. Forward-looking statements are statements other than historical fact and may include statements that address future operating, financial or business performance or Auris Medical's strategies or expectations. In some cases, you can identify these statements by forward-looking words such as "may," "might," "will," "should," "expects," "plans," "anticipates," "believes," "estimates," "predicts," "projects," "potential," "outlook" or "continue," and other comparable terminology. Forward-looking statements are based on management's current expectations and beliefs and involve significant risks and uncertainties that could cause actual results, developments and business decisions to differ materially from those contemplated by these statements. These risks and uncertainties include, but are not limited to, the timing and conduct of clinical trials of Auris Medical's product candidates, the clinical utility of Auris Medical's product candidates, the timing or likelihood of regulatory filings and approvals, Auris Medical's intellectual property position and Auris Medical's financial position. These risks and uncertainties also include, but are not limited to, those described under the caption "Risk Factors" in Auris Medical's prospectus relating to its Registration Statement on Form F-1, as amended, and future filings with the Securities and Exchange Commission. Forward-looking statements speak only as of the date they are made, and Auris Medical does not undertake any obligation to update them in light of new information, future developments or otherwise, except as may be required under applicable law. All forward-looking statements are qualified in their entirety by this cautionary statement.

Auris Medical Holding AG
Condensed Consolidated Interim Statement of Profit or Loss and Other Comprehensive Income (unaudited)
(in CHF thousands, except per share data)

	Three Months Ended June 30,		Six Months Ended June 30,	
	2014	**2013**	**2014**	**2013**
Research and development expenses	(4,201)	(4,411)	(8,350)	(6,384)
General and administrative expenses	(845)	(462)	(2,554)	(732)
Operating loss	(5,046)	(4,873)	(10,904)	(7,117)
Finance expense	(17)	(1)	(68)	(11)
Finance income	71	27	121	51
Loss before tax	(4,991)	(4,846)	(10,851)	(7,077)
Net loss attributable to owners of the Company	(4,991)	(4,846)	(10,851)	(7,077)
Other comprehensive income:				
Items that will never be reclassified to profit or loss				
Remeasurements of defined benefits liability	(407)	49	(424)	92
Items that are or may reclassified to profit or loss				
Foreign currency translation differences	(3)	4	3	(26)
Other comprehensive income	(410)	53	(421)	66
Total comprehensive loss attributable to owners of the Company	(5,401)	(4,793)	(11,272)	(7,011)
Loss per share, basic and diluted	(0.27)	(0.33)	(0.59)	(0.52)
Weighted average common shares outstanding, basic and diluted	*18,841,275*	*15,651,157*	*18,448,274*	*13,616,303*
Currency rate CHF / USD	*0.8888*	*0.9425*	*0.8907*	*0.9362*

Auris Medical Holding AG
Condensed Consolidated Interim Statement of Financial Position (unaudited)
(in CHF thousands)

	June 30, 2014	December 31, 2013
Assets		
Non-current assets		
Property and equipment	215	196
Intangible assets	1,483	1,483
Total non-current assets	1,734	1,678
Current assets		
Current financial assets and other receivables	694	525
Prepayments	714	183
Cash and cash equivalents	12,131	23,866
Total current assets	13,539	24,574
Total assets	15,273	26,252
Equity and Liabilities		
Equity		
Share capital	7,537	6,487
Share premium	48,233	35,608
Foreign currency translation reserve	57	54
Accumulated deficit	(44,295)	(33,116)
Total shareholders' equity attributable to owners of the Company	11,530	9,034
Non-current liabilities		
Employee benefits	745	328
Deferred tax liabilities	328	328
Total non-current liabilities	1,073	656
Current liabilities		
Convertible loans	-	13,711
Trade and other payables	1,105	954
Accrued expenses	1,564	1,897
Total current liabilities	2,669	16,562
Total liabilities	3,742	17,219
Total equity and liabilities	15,273	26,252
Currency rate CHF / USD	*0.8867*	*0.8894*

Contact:
Dr. Thomas Meyer, Chairman and CEO, +41 41 729 71 94, ear@aurismedical.com

Auris Medical
Cochlear therapies

Innovative Treatments for Inner Ear Disorders

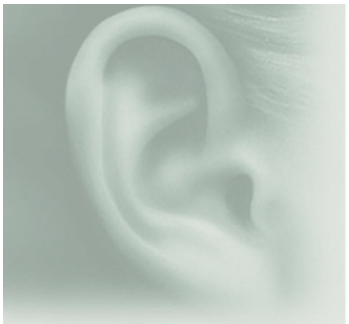
Second Quarter 2014 Earnings and Business Update

September 15, 2014

Forward Looking Statements / Safe Harbor

This presentation and the accompanying oral commentary contain "forward-looking" statements that involve substantial risks and uncertainties. All statements other than statements of historical facts contained in this presentation and the accompanying oral commentary, including statements regarding our future financial condition, business strategy and plans and objectives of management for future operations, are forward looking statements. In some cases, you can identify forward-looking statements by terminology such as "believe," "will," "may," "estimate," "continue," "anticipate," "intend," "should," "plan," "might," "approximately," "expect," "predict," "could," "potentially" or the negative of these terms or other similar expressions. Forward looking statements appear in a number of places throughout this presentation and the accompanying oral commentary and include statements regarding our intentions, beliefs, projections, outlook, analyses or current expectations concerning, among other things, our ongoing and planned preclinical development and clinical trials, the timing of and our ability to make regulatory filings and obtain and maintain regulatory approvals for our product candidates AM-101 and AM-111, our intellectual property position, our ability to develop commercial functions, expectations regarding clinical trial data, our results of operations, cash needs, spending of the proceeds from this offering, financial condition, liquidity, prospects, growth and strategies, the industry in which we operate and the trends that may affect the industry or us.

Forward-looking statements involve known and unknown risks, uncertainties, assumptions and other factors that may cause our actual results, performance or achievements to be materially different from any future results, performance or achievements expressed or implied by the forward-looking statements. Forward-looking statements represent our management's beliefs and assumptions only as of the date of this presentation. Except as required by law, we assume no obligation to update these forward-looking statements publicly, or to update the reasons why actual results could differ materially from those anticipated in the forward-looking statements, even if new information becomes available in the future.



Business Update

Auris Medical
Cochlear therapies

Innovative Treatments for Inner Ear Disorders

AM-101 Phase 3 Program Update

- AM-101 Phase 3 clinical program in acute inner ear tinnitus remains on track
- Ramp-up of TACTT trial sites progressing
  - Approximately 100 sites initiated by the end of August 2014 (target 140)
  - Poland and Czech Republic getting started
- As expected, Stratum B (post-acute tinnitus) in European TACTT3 trial enrolling faster than Stratum A (acute tinnitus)
  - Stratum B futility analysis early 2015 at midpoint of enrolment
- AMPACT open-label trials
  - First patients rolled over in April and May 2014 to receive first treatment cycle
  - First patients rolled over in July and August 2014 to receive second treatment cycle

Tinnitus Loudness Responder Definition

- Primary efficacy endpoint in Phase 3 = improvement of subjective loudness from baseline to the last follow-up visit at Day 84 on a 0-10 numerical rating scale

Tinnitus loudness on numerical rating scale "at its loudest" (TLQ $NRS_{Loudest}$)

On a scale from 0 to 10, where 0 represents no tinnitus and 10 represents extremely loud tinnitus, what one number best describes your tinnitus at its loudest **in the last 24 hours (including right now)**?

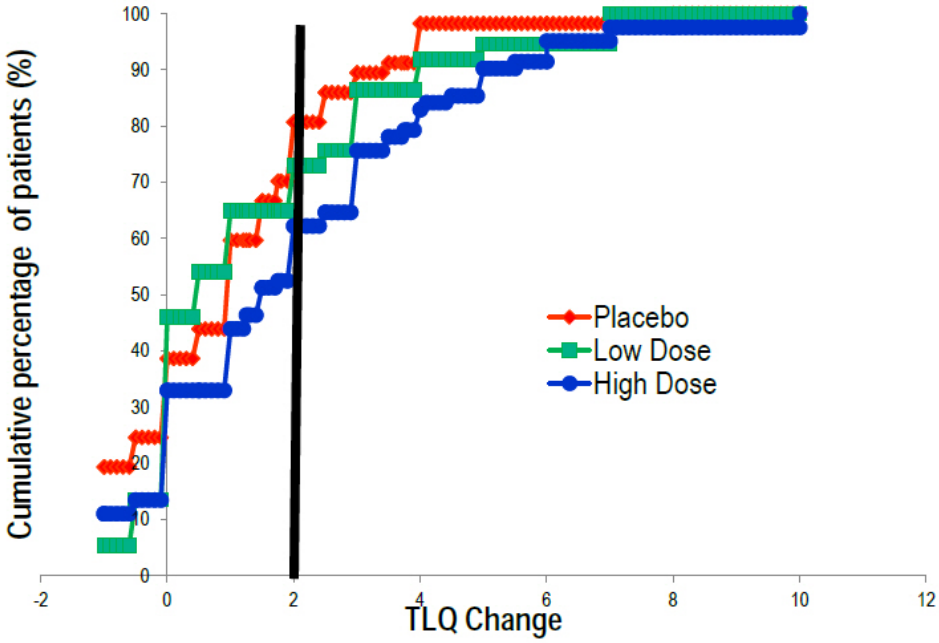
0	1	2	3	4	5	6	7	8	9	10
no tinnitus heard										extremely loud tinnitus

- What represents a clinically meaningful reduction in tinnitus loudness / how is a responder defined?
- Data analyzed from 200 TACTT0 & TACTT1 patients (AAT or OM)
- Mean tinnitus loudness at baseline = 5.2 points (SD 2.1)

Responder Data Presented at ITS 2014

- Results presented at XI. International Tinnitus Seminar in Berlin, Germany, May 21-24, 2014
- Anchor-based approach suggesting 2 point reduction as clinically meaningful
  - Pattern of TLQ response compared to patient global impression of change (PGIC) in tinnitus severity
  - Patients reporting either “very much improved” or “much improved” tinnitus severity reported change scores of at least 2 points, with those rating their change as “somewhat improved” reported less than a 2-point change
  - Patients endorsing the largest amount of change on the global rating also reported the largest amount of improvement on the TLQ
  - Pattern of response very similar for uni- and bilateral cases
- Supported by distribution-based approach:
  - One-half standard deviation of the baseline scores approach
  - Clinically meaningful = 1.1 points

Cumulative Distribution Function for Δ TLQ



- For all points > 0, the curve for AM-101 0.81 mg/mL is shifted to the right of the others
- At 2 points change in TLQ, AM-101 0.81 mg/mL clearly separates from placebo and AM-101 0.27 mg/mL, providing further support for responder definition

Auris Medical at the AAO Annual Meeting

- Annual Meeting of the American Association for Otolaryngology – Head and Neck Surgery
- Orlando FL, September 21-24
- World's largest gathering of ENTs
- Auris Medical hosting satellite symposium September 23

AM-101 Primary Market Research

- 53 US ENT doctors surveyed[1)]
  - 41 general ENTs
  - 12 otologists
- Seeing an average of 43.5 tinnitus patients in an average month
- 37.7% of their tinnitus patients seek treatment during the acute stage (up to 3 months from onset)
- Consistent with survey by Hall et al., 2011

- 73.6% of respondents expect their monthly tinnitus patient volume to increase if an approved i.t. treatment of acute tinnitus were available
- 42.6% of their tinnitus patients considered as candidates for a product with the profile of AM-101
- Rating
  - Tolerability 7.5
  - Efficacy 6.9

  (1=not at all attractive, 10=extremely attractive)

1) Online survey conducted by MedaCorp, Inc. in April 2014

AM-111 Program Update

- Preparations for the AM-111 late stage clinical program in ASHNL progressing
- Pivotal HEALOS Phase 3 trial planned in ISSNHL
  - 240 patients with severe to profound hearing loss
  - AM-111 0.4 or 0.8 mg/mL or placebo
  - Up to 72 hours from onset
  - Europe and Asia
- Protocol Assistance from European Medicines Agency (EMA)
- Pre-IND meeting with the FDA scheduled for second half of September 2014
- Based on FDA feedback, design of the late-stage AM-111 clinical program to be finalized and communicated

AM-111 Primary Market Research

- 53 US ENT doctors surveyed[1)]
  - 41 general ENTs
  - 12 otologists
- Seeing an average of 11.2 patients with ISSNHL and 6.3 patients with AAT in an average month
- 39.9% show up for treatment within first 3 days
- 43.3% of their ASNHL patients have severe to profound hearing loss

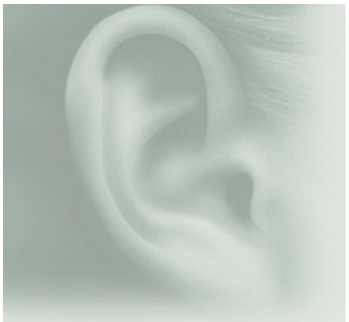
- 64.2% of respondents expect their monthly ASNHL patient volume to increase if an approved i.t. treatment of ASNHL were available
- 59.9% of their ASNHL patients considered as candidates for product with the profile of AM-111
- Rating
  - Tolerability 7.8
  - Efficacy 7.2

  (1=not at all attractive, 10=extremely attractive)

[1)] Online survey conducted by MedaCorp, Inc. in April 2014

Intellectual Property Update

- Additional patent applications filed in US relating to the use of inhibitors of the c-Jun N-Terminal Kinase (JNK) such as AM-111
  - for the treatment of tinnitus
  - for the treatment of Menière's Disease
- Continuation to earlier US application filed relating to the use of polymer-based formulations in the treatment of disorders of the middle and inner ear
  - Allowed by European Patent Office, covering compositions containing Ketamine or derivatives and biocompatible hyaluronic acid gel for treatment of inner ear disorders
  - Priority date = September 28, 2005



Financial Update

Auris Medical
Cochlear therapies

Innovative Treatments for Inner Ear Disorders

First Half 2014 Financial Highlights

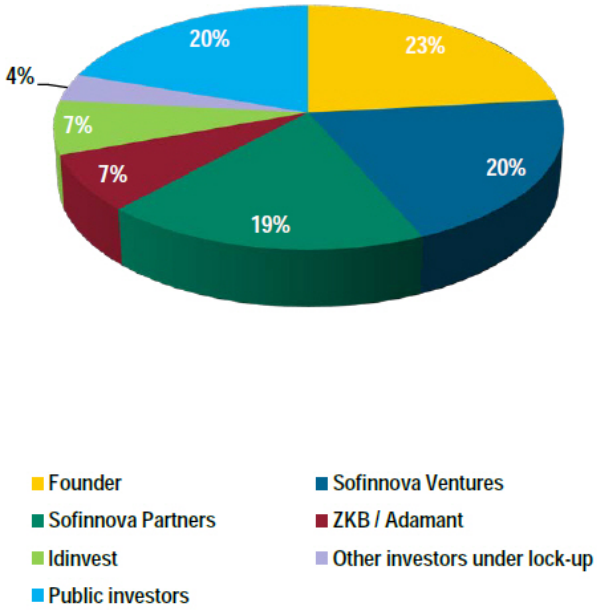
- Cash and cash equivalents as of June 30, 2014, were CHF 12.1 m compared to CHF 18.5 m at March 31, 2014 and CHF 23.9 m at December 31, 2013
- Research and development expenses increased from CHF 6.4 m in the six months ended June 30, 2013 to CHF 8.4 m in the six months ended June 30, 2014.
  - Progression of our AM-101 Phase 3 clinical development program
  - Preparatory activities related to the late-stage AM-111 clinical program
- General and administrative expenses increased from CHF 0.7 m in the six months ended June 30, 2013 to CHF 2.6 m in the six months ended June 30, 2014
  - Increase primarily related to higher legal and auditing expenses in relation to the IPO preparations
- Net loss for the first half 2014 was CHF 10.9 m compared with CHF 7.0 m in the same period in 2013.

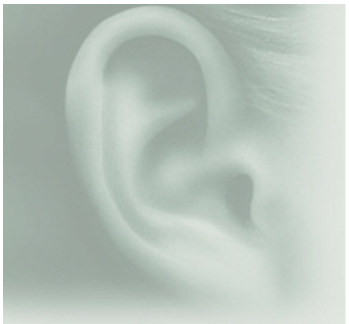
Financial Overview

Financial Position

- Cash at end of H1 2014 = $13.7 m
- $60.7 m raised in IPO in early August 2014 ($56.4 m after underwriting discount)
  - 10,113,235 common shares sold at $6 per share (incl. 713,235 shares under over-allotment option)
  - Trading on the NASDAQ Global Market started August 6, 2014
- Means to fund
  - AM-101 clinical program beyond Phase 3 read-out
  - Advance further projects, incl. AM-111
- Runway mid 2016

Shareholder Structure





Questions & Answers

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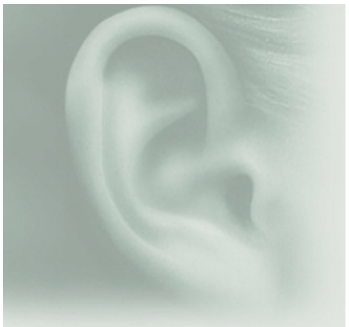

Take care of your ears!

Auris Medical
Cochlear therapies

Innovative Treatments for Inner Ear Disorders