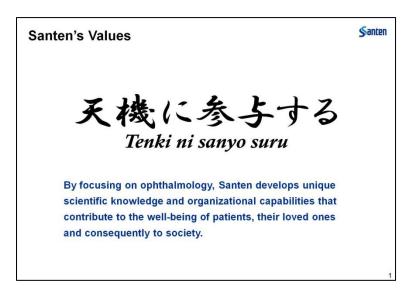
Speaker Remarks August 1, 2017 (summary)

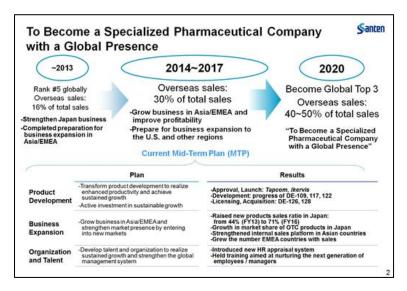
Santen's Values, Long Term Strategic Vision and MTP

Slide1: Santen's Values



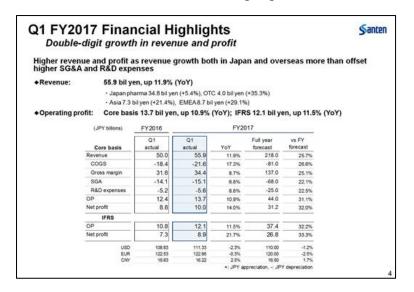
This is Koshiji speaking. I will use the slides. Please turn to Slide 1, this is our basic values.

Slide 2: To Become a Specialized Pharmaceutical Company with a Global Presence



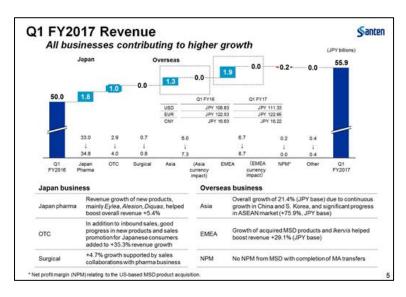
Slide 2, please. By 2020, we want to become a specialized pharmaceutical company with a global presence. This is our long-term vision. So, in 2014 to 2017, we are working on our medium-term plan to achieve that goal. The main initiatives are building on the growth of the business in Asia, EMEA and improving profitability and preparing for business expansion to the U.S. and other regions. This is the basis of our medium-term plan. 2017 is the final year, so let me explain the first quarter of this final year in our current medium-term plan.

Slide 4: Q1 FY2017 Financial Highlights



Slide 4, please. This is the outline of the first quarter results. Both revenue and operating profit grew by double-digits. As you can see here, both in Japan and overseas, we grew our revenues strongly. The overseas sales ratio is now 29%. Now the tax rate in the first quarter was 27%. So, this is down year-on-year.

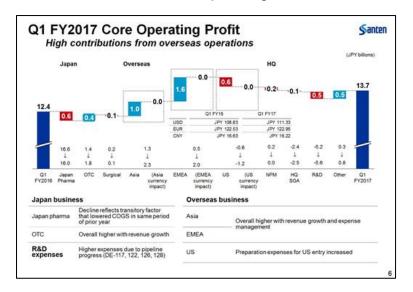
Slide5: Q1 FY2017 Revenue



Slide 5, please. This shows the trend of our revenue on bridge chart in comparison with previous year. As I said earlier, in Japan we had new pharma products, and inbound and new product in OTC, these were the drivers of growth. Regarding overseas, that is, Asia and EMEA: Asia grew by 20% and EMEA grew by 29%. This is yen denominated, but it is similar on a local-currency basis.

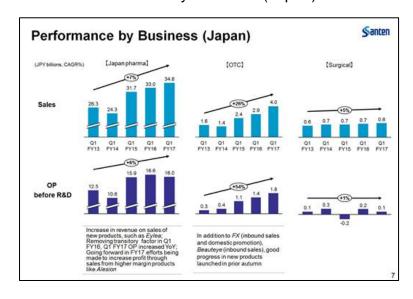
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Slide 6: Q1 FY2017 Core Operating Profit

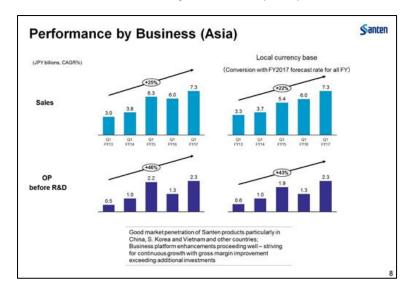


Now Slide 6, please. This is the comparison of our operating profit with the previous year. The bridge chart graph is the comparison with the previous year, and on the bottom half you can see the absolute operating profit. The upper row of figures is the previous year, and the lower row is this year. Japan pharma is the big component, but in terms of growth, overseas grew very strongly. Additionally, I would like to add that in the Japan pharma business, it reduced by ¥600 million. The operating profit declined on the accounting basis. But compared to the previous year i.e. because of the consumption tax hike, there was an adjustment of the procurement cost, which was a one-off factor. And because of that, the cost was lower last year by ¥700 million. So, against this high comparator, the profit in the first quarter of 2017 seems lower. But in real terms, both sales and profit increased.

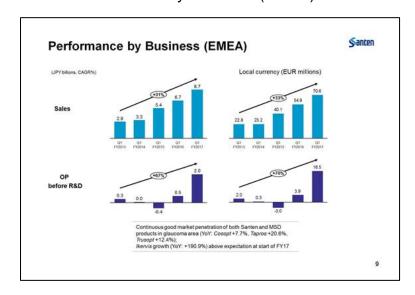
Slide 7: Performance by Business (Japan)



Slide 8: Performance by Business (Asia)



Slide9: Performance by Business (EMEA)



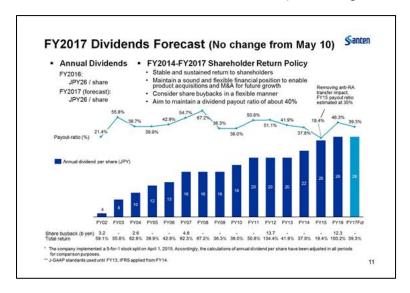
Slides 7, 8 and 9 show performance by business at each region.

Slide 10: FY2017 P&L Forecast (No change from May 10)

FY20	17 P&L I	Forecas	st (No cha	nge from	May 10)	Santi	
Core	Revenue: Growth	orecast in all busi	nesses, particularly	overseas			
basis	forecast to increase Investments in g	rating profit: Increased spending on future growth (listed below), while also strengthening cost control systems. OP is					
IFRS	Amortization on intangible assets associated with products will change substantially; non-recurring items; revenue and core operating profit to grow in proportion						
		(JPY billions)	FY2016 FY2017				
	Co	re basis	Actual	Forecast	YoY		
	Re	venue	199.1	218.0	9.5%		
		ogs	-75.0	-81.0	8.1%		
	5	GA	-61.7	-68.0	10.3%		
	F	&D expenses	-22.8	-25.0	9.7%		
	Op	erating profit	39.7	44.0	10.9%		
	Ne	t profit	28.7	31.2	8.8%		
		ROE	11.2%	12.3%	1.1pt		
	IFF	RS					
	Op	erating profit	32.5	37.4	15.2%		
	Ne	t profit	23.1	26.8	16.2%		
	-	ROE	9.0%	10.6%	1.6pt		
	_	USD	108.64	110.00			
		EUR	118.96	120.00			

Let's go to Slide number 10. This is our forecast which as I mentioned, has been progressing as planned. So, there has been no change from the forecast we announced on May 10.

Slide 11: FY2017 Dividends Forecast (No change from May 10)



Slide 11, this is about the dividend. Again, there has been no change made since the previous time. That's all from me. Thank you.

Slide 19: Future Development and Regulatory Milestones

	Development status*	Indication	Development region: milestone*	
DE-117	P2b/3	Glaucoma/ ocular hypertension	Japan: Q2/Q3 FY17 filing	
DE-126	<u>P2b</u>	Glaucoma/ ocular hypertension	US/Japan: Jan~Jun 2018 P2b completion	
DE-128 (MicroShunt)	P2/3	Glaucoma/ ocular hypertension	US: calendar 2018~2019 P2/3 completion, calendar 2020~2021 launch	
DE-109	Filing Accepted	Uveitis	US: Dec 24, 2017 PDUFA date; Jan~Jun 2018 launch	
(IVT sirolimus)	P3	Oveids	EU: 2 nd half of FY17 re-filing	
DE-122	<u>P2a</u>	WetAMD	US: <u>Jan~Jun 2019 P2a completion**</u> , FY17 P1/2 completion	

I'm Naveed Shams and I will start on Slide 19.

And I'll be starting with our product, the DE-117, which is currently in some regions in Phase 3, Phase 2b/3 development. However in Japan, we have finished all the required studies. And as planned, we'll be filing by the third quarter of fiscal 2017. This will be treatment for glaucoma and ocular hypertension.

The second glaucoma IOP-lowering product that we have recently put into development is DE-126. This is a Phase 2 study in both U.S. and Japan. And we hope to complete the study by June or so, middle of next year.

Staying with the glaucoma theme, our flagship product is another one is DE-128, the MicroShunt device. The enrolment for the pivotal study, required for launch in the U.S., is going quite well. And so, we continue with our timeline of completion of the study in 2019.

DE-109 which is a non-infectious uveitis product was filed early in the year, and the review is going without any major issues. We expect our approval date to be the week of December 24.

Another product DE-122 is for wet AMD, and that study has started in the Philippines, and we expect completion of this Phase 2 study in around the middle of 2019.

I'm also delighted to tell you that our ciclosporin product called Verkazia was given a positive opinion by the EMA's CHMP recommending to the commission that it be approved for marketing in the European Union.

We have also started a line extension of our allergy product, Alesion which is epinastine hydrochloride, DE-114A. And, it started Phase 3, a couple of months ago in May 2017. This is to actively manage the allergy franchise which is very important product for Santen.

Thank you very much, that's all for my side.