Case 1:16-cv-01221-LPS Document 161-1 Filed 07/06/20 Page 1 of 7 PageID #: 1352

Exhibit A

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		Page 1			
1	UNITED STATES DISTRICT COURT				
2	FOR THE DISTRICT OF DELAWARE				
3					
4	BAYER HEALTHCARE LLC and BAYER)				
5	HEALTHCARE PHARMACEUTICALS, INC.,) Plaintiff,) Case No.				
6	v.) 16-1221 (LPS) - USDC-DDE				
7	TEVA PHARMACEUTICALS USA, INC.,) APOTEX, CORP. AND APOTEX INC.,)				
8	Defendants.				
9	/				
10					
11	CONFIDENTIAL				
12					
13					
14	DEPOSITION OF ALLAN MYERSON, Ph.D.				
15	Washington, D.C.				
16	October 21, 2019				
17					
18					
19					
20					
21					
22	REPORTED BY: Tina Alfaro, RPR, CRR, RMR				

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Case 1:16-cv-01221-LPS Document 161-1 Filed 07/06/20 Page 3 of 7 PageID #: 1354

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Page 82 11:11:14 1 that. 11:11:18 2 Q. Well, let's just kind of break that up. 11:11:20 3 So there are four stages in the example, correct? 11:11:24 4 A. Correct. 11:11:28 And looking at stage 3, which is on 5 Q. 11:11:32 column 14, what amount of impurities do you 6 11:11:37 7 achieve? 11:11:39 8 Α. It doesn't say. 11:11:45 9 Certainly doesn't tell you whether Q. 11:11:47 10 0.0001 percent was ever achieved, correct? 11:11:53 11 A. It doesn't say, but, of course, a POSA 11:11:56 12 could perform this example and do the analysis and 11:11:58 13 they would know. 11:12:03 14 Q. Stage 4 also measures the amount of 11:12:05 15 impurities, correct? 11:12:06 A. Correct. 16 11:12:09 17 Q. And it does not tell you whether 11:12:11 0.0001 percent was ever achieved, correct? 18 11:12:17 A. The data's not in there, that's correct. 19 11:12:21 20 The POSA would have to perform example 4 and 11:12:23 determine if it was achieved. 21 11:12:25 22 Q. To achieve 0.001 percent can you show me

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Case 1:16-cv-01221-LPS Document 161-1 Filed 07/06/20 Page 4 of 7 PageID #: 1355

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Allan Myerson, Ph.D. - October 21, 2019

Page 83

			rage 00
11:12:34	1	in the patent where it says what changes need to be	
11:12:37	2	made to the basic synthetic pathway?	
11:12:41	3	A. I think you mean 0.0001 percent. You left	
11:12:45	4	a zero out again.	
11:12:47	5	Q. Yes.	
11:12:48	6	A. Yeah. If you want to make this simple for	
11:12:51	7	yourself, you could just say 1 PPM and a hundred	
11:12:54	8	PPM if that's easier to do.	
11:12:57	9	Q. Let's do that. 1 PPM, just so we're	
11:13:01	10	clear, is let's try this again	
11:13:05	11	0.0001 percent. Fair enough?	
11:13:10	12	A. I think you left a zero out again.	
11:13:13	13	THE REPORTER: No, he didn't.	
11:13:15	14	THE WITNESS: He didn't that time? Good.	
11:13:16	15	Okay.	
11:13:17	16	BY MR. MALIK:	
11:13:17	17	Q. 1 PPM, I actually do like that much	
11:13:20	18	better.	
11:13:21	19	The patent doesn't show you what specific	
11:13:23	20	changes need to be made to achieve 1 PPM versus a	
11:13:26	21	hundred PPM, does it, to the basic synthetic	
11:13:28	22	pathway?	

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Case 1:16-cv-01221-LPS Document 161-1 Filed 07/06/20 Page 5 of 7 PageID #: 1356

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			Page 8	34
11:13:29	1	A. Say that again at the end. I'm sorry.		
11:13:32	2	Q. Sure. Is there any strike that. Let		
11:13:37	3	me see if I can lay some foundation.		
11:13:40				
	4	In paragraph 87 of your report you say		
11:13:43	5	that to achieve the levels would require		
11:13:46	6	optimization involving a variety of variables,		
11:13:48	7	including the basic synthetic pathway, reaction		
11:13:52	8	conditions, and you go on, correct?		
11:13:54	9	A. Right.		
11:13:54	<mark>10</mark>	Q. Okay. So that's the foundation for my		
11:13:56	11	question. The patent doesn't show what specific		
11:14:06	<mark>12</mark>	changes need to be made to the basic synthetic		
11:14:09	<mark>13</mark>	pathway to achieve 1 PPM versus 100 PPM's, correct?		
<mark>11:14:15</mark>	<mark>14</mark>	MR. BOWERS: Object to the form of the		
<mark>11:14:16</mark>	<mark>15</mark>	question.		
11:14:18	<mark>16</mark>	A. Okay. I think I understand. The patent		
11:14:22	<mark>17</mark>	itself teaches the optimized synthetic procedure to		
11:14:29	<mark>18</mark>	make regorafenib with the desired levels of		
11:14:33	<mark>19</mark>	impurities. Practicing the examples should allow		
11:14:37	20	you to do so, but if you're asking me it doesn't		
11:14:43	<mark>21</mark>	it doesn't say if I do it one way I'll get a		
11:14:46	22	hundred PPM and if I do it this way I'll make 1		

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