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NYSCEF DOC. NO. 1706

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EXHIBIT N

1	UNITED STATES DISTRICT COURT
2	CENTRAL DISTRICT OF CALIFORNIA - WESTERN DIVISION
3	HONORABLE S. JAMES OTERO, U.S. DISTRICT JUDGE
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6	JUNO THERAPEUTICS, INC., et al.,)
7	Plaintiffs,)
8	vs.) 2:17-CV-7639-SJO
9	KITE PHARMA, INC.,
10	Defendant.))
11	
12	REPORTER'S TRANSCRIPT OF PROCEEDINGS
13	JURY TRIAL VOLUME IV
14	
15	Los Angeles, California
16	Friday, December 6, 2019
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22	AMY DIAZ, RPR, CRR, FCRR Federal Official Reporter
23	350 West 1st Street, #4455 Los Angeles, CA 90012
24	LOS ANGELES, CA JUUIZ
25	Please order court transcripts here: www.amydiazfedreporter.com

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1	MR. HEINRICH: All right. PX78, 155, 156, 240
2	MR. WEINBERGER: 240 has an objection, Your Honor.
3	THE COURT: Okay. 240 is not in.
4	MR. HEINRICH: So we may have different lists, so
5	I'll do the rest one at a time.
6	THE COURT: So any so any objection to the 78,
7	155 and 156?
8	MR. WEINBERGER: No, Your Honor.
9	THE COURT: Okay. Those exhibits are received.
10	(Exhibits PX78, PX155, and PX156 received.)
11	DIRECT EXAMINATION
12	BY MR. HEINRICH:
13	Q. Good morning, Dr. Sullivan.
14	A. Good morning.
15	Q. Can you please introduce yourself to the jury.
16	A. Good morning. I am Ryan Sullivan. I work as an
17	economist. And I serve as president of Intensity.
18	Q. And why are you here?
19	A. To share my expert analysis and opinions relating to
20	money damages that were incurred by Juno as a result of the
21	alleged infringement.
22	Q. And have you prepared anything to help illustrate your
23	testimony today?
24	A. Yes. I prepared a set of demonstratives.
25	Q. All right. So can you please summarize your educational

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background briefly? 1 2 A. Yes. I have a bachelor's degree, a master's degree and a 3 Ph.D. They are all in economics and all from the University of California in San Diego. 4 Q. Did you continue working with the University of San 5 Diego -- University of California San Diego? 6 7 A. Um, I did. I served as an invited member of the 8 Economics Leadership Council for the Department of Economics at UCSD, wherein I advised the faculty at UCSD on the 9 10 practice of economics and private industry. Q. Have you published any work in economics or economic 11 12 damages? 13 A. Yes, I have. I have a number of publications in peer-reviewed journals, including the Journal of Finance, the 14 Journal of Econometrics, and the International Journal of 15 Forecasting. I have also published my research in les 16 17 Nouvelles, which is the journal for the Licensing Executives 18 Society as well as other papers involving intellectual 19 property. 20 Q. And how long have you been providing professional 21 economic services? A. Since April 1992. So it has been just over 27 years now. 2.2 23 Q. You mentioned you are president of Intensity. What is 24 Intensity? A. We are an economics and data science firm with about 20 25

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Ph.D. economists and business analysts. We work with companies to help them solve problems and strategic decision making in the marketplace. Q. What were you asked to do in this case? A. I was asked to provide my expert research and analysis to the Court to provide quidance in determining what the appropriate reasonable royalty would be for the use of the '190 Sadelain patent by Kite. Q. And as part of that analysis, do you have to make any assumptions in connection with that? The standard assumption for calculating damages is to assume that the patent is both valid and infringed. Q. Now, what materials have you considered in forming your opinions? And let's go to slide 3. A. A great number of documents and data and information, so this includes documents that were produced by the companies to this litigation as well as filings that they have made, for example, with the Securities and Exchange Commission. I have looked at financial data on sales and profitability, market research reports and analysis that has been performed. I've also reviewed extensive testimony from witnesses within the litigation. I've held interviews with other experts. And I have also analyzed and utilized company financial models. O. Now, what is a reasonable royalty?

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SULLIVAN - DIRECT A. It is a form of damages in patent cases. As an economist, I perform my work in the context of the law. the statute provides for damages adequate to compensate for the infringement but in no event less than a reasonable royalty. Q. And how do you evaluate a reasonable royalty? A. Yeah. A standard way to do so is through what is known as a hypothetical negotiation. This would be a negotiation that would happen between the licensor, parties with a patent, and a licensee, parties that are seeking to obtain a license or permission to use that patent. It's hypothetical in a couple of ways. One, the negotiation did not actually happen. So we have to reconstruct what that negotiation would look like.

One, the negotiation did not actually happen. So we have to reconstruct what that negotiation would look like.

Also, we assume that the patent is valid and infringed, which is different than a lot of real world negotiations. And we also look at all of the information that is available and recognize that the parties to this hypothetical negotiation would have access to the whole collection of information.

- Q. Now, who would have participated in the hypothetical negotiation that you analyzed?
- A. Well, in the hypothetical negotiation in this case, that would have occurred on October 18th, 2017, or right before that. And that's the date upon which Kite received FDA approval to begin selling YESCARTA.

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And so at that point in time, Kite Pharma would be the entity that would be seeking a license. And just two weeks prior to that point in time, they had been acquired by Gilead. So Gilead would also be very interested in the outcome of the negotiation as they would be seeking and wanting to launch YESCARTA into the marketplace.

On the other side would be Memorial Sloan Kettering, Sloan Kettering, and Juno as the ones that are interested in the rights to the '190 Sadelain patent.

- Q. Now, you mentioned that some of the materials you considered were company financial models. What role did they play in your hypothetical negotiation analysis?
- A. These are Excel spreadsheets and financials that are put together by the companies in the ordinary course of business, not for purposes of this litigation, but for making business decisions, whether it be prioritizing products or evaluating acquisitions.

And these are really informative in this case for the hypothetical negotiation because it gives us direct insight into how the parties would be thinking at the hypothetical negotiation. In other words, what their expectations would be at that point in time.

- Q. Are there factors that are typically considered in a reasonable royalty analysis? And this is slide 8.
- A. Yes. About 50 years ago there was a case, very

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well-known case involving Georgia Pacific, and the Court there set forth 15 different factors that may be informative for determining a reasonable royalty. And I considered and evaluated all 15 of these factors in the course of my work. Q. Did you find any of those factors particularly relevant to this case? A. Yes, I did. In particular, factors 9, 10, and 11, those inform upon the importance of the Sadelain patent. Factors 8, 12, and 13 inform upon the financial benefits to Kite of utilizing the patented technology. Factor 5 addresses the competitive relationship between Kite and Juno. And factors 1 and 2 inform upon license agreements and licensing for technology. Q. All right. Let's take a look at some of the evidence here. MR. HEINRICH: So I would like to move into evidence a redacted version of PX72 that just includes the cover page and a paragraph on PX72.81. THE COURT: Let me find it first. MR. WEINBERGER: Excuse me, Your Honor. We don't have the plaintiffs' binder here. I don't have the redacted version. THE COURT: I think they are trying to secure it for you. MR. HEINRICH: We do have the version on our

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1	electronic equipment. Can I pull up the unobjected-to
2	paragraph?
3	THE COURT: Why don't you confer with counsel and
4	make sure it's agreeable.
5	MR. HEINRICH: May I approach with redacted copies?
6	THE COURT: Yes, please. If you can hand it to
7	Mr. Cruz.
8	MR. WEINBERGER: Subject to previous objection we
9	made about this.
10	THE COURT: Yes. The objections previously noted
11	will remain.
12	And so Exhibit 72, is this page 1? What is it?
13	MR. HEINRICH: Page 1 and redacted page 81 on the
14	second I apologize, that is double sided.
15	THE COURT: On the 72.81, the redacted, there is
16	nothing there.
17	MR. HEINRICH: There is the one paragraph that is
18	the paragraph that we wanted to use here.
19	THE COURT: Okay. Not on my copy, but go ahead.
20	There is no objection.
21	BY MR. HEINRICH:
22	Q. So what is Exhibit 72, Dr. Sullivan?
23	A. This is a form 10K that was filed by Gilead with the
24	Securities and Exchange Commission for the year ending 2017.
25	And a form 10K is an annual report that public companies file
	d .

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with the Securities and Exchange Commission to provide 1 2 detailed information about the company to the SEC as well as 3 to investors and shareholders and provides detailed information regarding the company. 4 Q. And what information did you find in this exhibit 5 regarding the value of the Sadelain patent to YESCARTA? 6 7 A. On page 81 there is information that I found to be very 8 informative in this regard. So first I'll just read a 9 relevant piece here, and then I can explain what it means. 10 So starting on the second line, "In October 2017, 11 upon FDA approval of YESCARTA for the treatment of adult 12 patients with relapsed or refractory DLBCL after two or more lines of systemic therapy, \$6,200 million of the purchased IP 13 R and D was reclassified as a finite lived intangible asset." 14 15 Q. Can you help translate that for us? 16 A. Yes. So what this is saying is that when Gilead acquired 17 Kite, that there was a portion of that purchase price that they identified as being the value attributable to IP R and D 18 of YESCARTA. IP R and D is intellectual property research 19 20 and development. So think of that as the technology in 21 YESCARTA. What they have done here is identified what the 2.2 value is of that item for YESCARTA, and that is the 23 24 \$6,200 million. Q. What is \$6,200 million? 25

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A. That is an odd way of saying \$6.2 billion. Most all of the numbers in the form 10K are rather large, so they are all treated as millions. Here, it is 6,200 million. We would more likely refer to that as just over \$6 billion.

And the way to think about value here, somewhat of a formal or structured way of thinking about it, is that value is based upon the expectation of future profits. That is what gives an asset value. And, naturally, the further out in time into the future that one is projecting, one has to account for that in the value.

In other words, there is two adjustments to kind of think about here. One is that you want to have your -- you know, money today is worth more than money received a year from now or five years from now. But there is also a little bit more uncertainty associated with the future. And what the value does is it looks at what is the present value today, taking all of that into account. And that is what Gilead determined and reported as being \$6.2 billion.

- Q. So from an economics standpoint, what is your understanding of how YESCARTA uses Dr. Sadelain's patented CAR?
- A. As I understand it, that patented CAR is what allows
 YESCARTA to be a living drug with a single dose of therapy to
 provide the efficacy and the therapeutic value of the
 product.

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Q. Now, have you seen any evidence from Kite about the role 1 2 of the Sadelain patent -- the Sadelain patented CAR in 3 YESCARTA? A. Yes, I have. 4 Q. And let's pull up our next slide. 5 And can you explain this evidence to the jury? 6 7 A. Yes. So this is testimony, and we heard this just a little bit earlier this morning, from Shawn Tomasello who was 8 the former chief commercial officer at Kite, and she 9 explained that it's fair to say that having a construct on 10 11 which to base your product is an important aspect from a 12 commercial point of view and that, without that CAR construct, there isn't anything. In other words, it's that 13 CAR construct that is fundamental to the product and 14 ultimately, as we'll see, it really is fundamental to all of 15 the business for CAR-T for Kite. 16 17 Q. Did any of Kite's real world activities shed light on the 18 value of the Sadelain patent to Kite at the time of the hypothetical negotiation? 19 20 A. Yes. There are several key items in this regard. 21 The first is that Kite attempted historically to 2.2 obtain a license to the '190 Sadelain patent, but they were 23 unable to do so. They did not secure a license that would 24 give them permission to use the '190 patent. Subsequent to that point in time, they then sought 25

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to invalidate the patent through what is known as an inter partes review process. That attempt to invalidate the patent was --MR. WEINBERGER: Objection, Your Honor, I think this is far afield. THE COURT: The objection is -- ask your next question. The objection is overruled. BY MR. HEINRICH: Q. Did you see any evidence that Kite tried anything else with respect to the Sadelain patent? There were multiple attempts by Kite to develop alternative constructs. There is the Kite 585 construct, which was an internally-developed construct. But ultimately they did not continue with that as they found it was not efficacious. There is another construct referred to as HU19. That construct they brought through to phase I clinical trials and then decided not to pursue that beyond phase I. So there were two other attempts, and neither one turned out to be successful for them and are focused in their CAR-T program solely on the Sadelain construct. Q. Well, do you have an understanding of the extent to which Kite's CAR-T program pipeline uses the Sadelain patented CAR? A. Yes. The ZUMA-1 trial, which resulted in YESCARTA for 3L DLBCL is one of the trials, and that's what we were

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discussing earlier that was valued by Gilead at \$6.2 billion. 1 2 But there are at least 10 other trials that are in 3 the pipeline for Kite that also are all using this very same Sadelain CAR construct. 4 Q. Now, are there other aspects or factors related to 5 YESCARTA that -- or -- that are important, that you 6 7 considered? A. There are other factors. And there are other 8 contributors to YESCARTA beyond just the construct itself. 9 10 For example, there's manufacturing processes, and 11 administration, and lymphodepletion. Yet, these other 12 factors do not drive the demand for the product. Because without the CAR construct, there isn't anything to 13 manufacture, there is no basis for lymphodepletion on a 14 15 patient. So it really is that the CAR construct is that which 16 17 is fundamental in the driver of demand for sales of YESCARTA. 18 Q. But did you take those other factors into consideration 19 in your analysis? 20 A. I did. And my analysis fully captures that value, and 21 separates out the value, contribution of the '190 patent relative and separate from all of the other contributions. 2.2 23 Q. So let's turn to the relationship between Juno and Kite. 24 How would you characterize that relationship? A. Juno and Kite are and have been direct competitors, and 25

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the evidence shows that they have considered each other to be 1 2 primary competitors. 3 Q. And what are some of the ways in which they have competed in the time period leading up to the hypothetical 4 negotiation? 5 A. On the next slide, I have a list of various items. There 6 7 are -- there has been competition, for example, for 8 technology rights, and in particular with regards to rights to the '190 patent. There has also been competition with 9 10 regards to funding. So obtaining investment, you know, both companies were seeking to go public and did have their IPOs 11 12 right around the same time. So they have been directly 13 competing in the space for investments. Also, with respect to employees, they have competed. 14 15 And competing for clinical trial sites. So the two entities have had a relationship that is directly competitive. 16 17 Q. Now, at the time of the hypothetical negotiation in 18 October of 2017, was there an expectation of future competition between Juno and Kite? 19 20 A. Yes. All of the documents produced by the entities in 21 these company financial models that I mentioned earlier 2.2 recognize that there would be overlap in a number of 23 different indications for the two entities, such that the 24 competition would continue.

Q. Now, was there an advantage to Kite in getting a license

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to the Sadelain patent at the specific time of the 1 2 hypothetical negotiation? 3 A. Yes. And, in fact, having a license upon launch in October 2017 provided Kite with a very significant 4 first-mover advantage. 5 Q. Let's turn to slide 20. What is a first-mover advantage? 6 7 A. This is an advantage that on net results in a company 8 that is first to providing their product into the marketplace with an advantage in terms of increased market share, 9 10 increased sales, being able to develop and shape the 11 marketplace so that there is increased physician loyalty and 12 loyalty from treatment centers, such that there can be a much 13 longer and entrenched advantage as a result of being first in 14 the marketplace. Q. Let's pull up Exhibit 29. It's in evidence. 15 16 And can you remind us what this is? 17 This is a presentation that provides the results of 18 research that was conducted by Kite internally, just a bit before the hypothetical negotiation, addressing the value of 19 20 a first-mover advantage. 21 Q. Let's turn to page 2. What is Kite saying here? 2.2 A. Here they are saying that based upon their research that 23 they did, that on average first movers achieve a higher 24 market share. 25 Q. And if we turn now to page 3.

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A. Here they are stating that based upon their research that the market share advantage of first movers at year 10 is on average 6 percentage points. And as an economist, this is really interesting, because one can think of a first mover having an advantage early on, and that perhaps that advantage would dissipate or start to diminish or go away perhaps quickly.

But the facts actually show that the advantage continues for a long period of time. And what they're saying here is that on average after 10 years, there is still a benefit of 6 percentage points at the market share.

- Q. And if we can turn to page 7.
- A. These are the conclusions of the study based upon their research that specifically to KTEC19, which is YESCARTA, that that first-mover advantage was expected to be between 5 and 10 percentage points continuing after even 10 years. And that part of the reason is that the treatment centers would gravitate towards one company's offerings across indications to be able to simplify the treatment process.
- Q. So let's pull up Exhibit 78. And while we pull that up, can you tell us whether you saw any evidence that Gilead considered that there was a first-mover advantage for YESCARTA?
 - A. Yes.

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Q. So let's turn to page 5 of Exhibit 78. Can you explain

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1 what we see here? 2 A. Yes. So this is a document that was prepared on the day 3 that Gilead announced to the marketplace that they are going to be acquiring Kite. And this is a document that was 4 circulated among the top executive team at Gilead to be able 5 to help answer questions that they might receive regarding 6 7 their decision to acquire Kite. 8 And what's interesting here is there is a question of why Kite instead of Juno. And what they are asking is, 9 10 why did Gilead decide to acquire Kite instead of Juno? And 11 the response that the company put together is because of the 12 first-mover advantage. That was the top reason that they 13 listed. Q. Did you see any other evidence from Gilead relating to a 14 15 first-mover advantage and its position on that? A. Yes, I did, there is what's referred to as earnings call 16 17 transcripts. So for public companies, they will hold 18 earnings calls with investors, and these calls are public. And there are transcripts made of what the executives 19 20 communicate into the marketplace regarding the company and 21 market developments. 2.2 MR. HEINRICH: So we move PX240 into evidence, which 23 is a Gilead 2017 earnings call transcript. 24

THE COURT: Any objection?

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MR. WEINBERGER: It's hearsay. It's not a Gilead

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1	document.
2	THE COURT: 240?
3	MR. HEINRICH: Yes.
4	THE COURT: And the objection is hearsay. The claim
5	is it's not a Gilead document?
6	MR. WEINBERGER: Correct, Your Honor.
7	THE COURT: Then lay a foundation, please. Lay a
8	foundation.
9	BY MR. HEINRICH:
10	Q. All right. Well, does the call transcript reflect
11	expectations of the parties at the time of the hypothetical
12	negotiation?
13	A. Yes. So during these calls, the executives and in
14	particular here, John Milligan, the chief executive officer
15	of Gilead, is in a public forum providing his input on the
16	marketplace and on Gilead, in this particular instance,
17	regarding a first-mover advantage associated with regards to
18	Kite. And Thomson Reuters is the entity that actually
19	transcribes the call and then makes the transcript available
20	to all of us out in the public.
21	Q. And as a damages expert, do you rely regularly on
22	earnings call transcripts such as this?
23	A. Yes. This is just part of our typical market research
2 4	that we would perform.
25	MR. HEINRICH: With that, we

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1	THE COURT: 240 is now received.
2	(Exhibit PX240 received.)
3	BY MR. HEINRICH:
4	Q. So if we pull up page 12. So can you tell us what so
5	first, who is John Milligan?
6	A. So the president, CEO, and director of Gilead.
7	Q. And what's reported here?
8	A. He states he states that he thinks the first mover is
9	very important with this area, because it allows them to
10	develop the rapport and relationship with the different
11	cancer centers. And so he thinks a first-mover advantage is
12	important in this area.
13	Q. Dr. Sullivan, were you here in court yesterday for Arie
14	Belldegrun's testimony?
15	A. Yes, I was.
16	Q. And did you hear his testimony about first mover not
17	being important to Kite?
18	A. I did.
19	MR. WEINBERGER: Objection, mischaracterizes his
20	testimony.
21	THE COURT: Overruled.
22	BY MR. HEINRICH:
23	Q. And what is your what is your opinion on that based on
24	all of the documents that you have reviewed in this case?
25	A. I found the testimony to be odd. All of the documents,

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all of the analysis, all of the research all points to the fact that there is indeed a first-mover advantage.

While certainly there may be certain aspects that aren't as beneficial, I think it's very clear that on net, overall, this was a very significant advantage for Kite.

- Q. Now, did you consider whether Juno would have anticipated how Kite's first-mover advantage would impact Juno's business?
- A. Yes. Evidence demonstrates and even economics and basic common sense that when Kite has the first-mover advantage and they receive those benefits, there is a reciprocal harm to Juno from being second or being behind Kite and further behind. Because if you can even think about it, if there are two products coming to market and suppose they are very similar, one of them gets there first, gets entrenched. The second one, when they are coming in, they no longer can be equal or similar. They have to be better in order to overcome those hurdles. So it puts them at a significant disadvantage. It slows down the process, slows down the launch, and causes them to have lower market share and thus less revenue and profitability over time.
- Q. So let's go back to the slides and talk next about your approach in evaluating reasonable royalties. At the hypothetical negotiation, what type of royalty structure would the parties agree to?

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A. The appropriate structure that the parties would agree to is a reasonable royalty that would have two components to it. One, an upfront payment, and two, a running royalty rate that would be applied to sales of YESCARTA. The second component sometimes is casually referred to as royalties, yet it's the two components combined that provides for the total reasonable royalty. Q. And why do you believe the parties would have agreed to this structure with both an upfront and a running royalty rate on sales? A. There's three good reasons for this. One, nearly all of the agreements that have been entered into by Kite and by Juno have this type of structure. Second, throughout the biotech industry, this structure is very common and predominates. And third, it's -- I label the title here agreed license structure, and that's because both Kite's economics expert, Dr. Rao, and I, we both agree that the reasonable royalty would be of this structure and have both an upfront payment and a running royalty on sales. Q. Dr. Sullivan, what methodology did you use to quantify these two components of your suggested royalty? A. I used a standard approach that is referred to as the

market approach. Sometimes it's also referred to as a

comparable license approach.

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And that's wherein I identify a comparable license agreement, and then make adjustments to that to have it be consistent and reflect what would be the outcome of the hypothetical negotiation. Q. Did you make adjustments to both components of this license? Both the upfront payment and the running royalty in this case requires adjustment to be able to reflect the differences between the agreement on the one hand and the hypothetical negotiation on the other. And I'll explain that in more detail. Q. Let's turn to slide 23. So how did you apply the market approach methodology? A. So given all the context, the research that I did, and the analysis that I've already been discussing, I determined that the appropriate agreement to start with is the 2013 exclusive license agreement between Memorial Sloan Kettering and Juno. And then made adjustments to that to reflect what would actually have occurred from the hypothetical negotiation, because there's very significant differences between that agreement being entered into in 2013, and the hypothetical negotiation in 2017 with Kite. Q. So what are those major differences between the Juno/MSK

2013 exclusive license and the hypothetical negotiation?

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A. So in 2013, that agreement was between collaborators, between Sloan Kettering and Juno.

It was also at a time when the CAR-T marketplace was in very early stages. In contrast, the hypothetical negotiation doesn't occur until October 2017.

During this time, although Kite is undertaking development of YESCARTA and seeking to get it launched, that period of time, that is not considered infringement. Rather, the infringement begins in October 2017 upon FDA approval and the launch of YESCARTA.

At that point in time, the market is far more developed, there is a fully commercialized product that is being launched in the marketplace, and the parties to the hypothetical negotiation are primary competitors.

So there's significant differences between the agreement on the one hand and the hypothetical negotiation that we have to account for, yet that initial agreement from 2013 between Sloan Kettering and Juno is the right place to begin, because it's for the very technology that's at issue in this case. And there really is no dispute that that's the right place to begin.

Again, Kite's economics expert Dr. Rao and I both agree on that point.

Q. Now, how do you account for the differences you just identified between the MSK/Juno 2013 agreement and the

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hypothetical negotiation?

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A. I take it in two steps. And that's because there is information that helps us get a part of that difference accounted for, and then another step that addresses the full difference.

So you'll see on the next slide or as we progress that there was an agreement in 2013, separate agreement between St. Jude and Juno that licensed a different patent.

And then as part of a settlement in 2015, Juno then licensed that patent to Novartis.

And so I used those two agreements to make one adjustment, but that only gets us part of the way there. Because that only gets us to 2015. That's still relatively early stage, we don't have a launched product. And secondly, that's with Novartis, not with Kite. And Novartis was a lesser competitor. And I'll be able to quantify that for you. And so I make a second adjustment to get us to Kite and Juno in 2017 at the hypothetical negotiation.

THE COURT: Let's take the noon recess. Please return back to the courtroom at 12:30, and we'll continue with the trial. During your absence, do not discuss the case amongst yourself or with any other person, please.

THE CLERK: All rise, please.

(Thereupon, the jury retired from the courtroom.)

(Thereupon, there was a lunch recess.)

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1	THE COURT: We have everyone present on Juno vs.
2	Kite, I think everybody is present. Mr. Chu is not here.
3	Does he intend to be here?
4	MR. HEINRICH: He does, Your Honor. Let's wait a
5	minute.
6	THE CLERK: I'm going to line up the jury.
7	MR. DANE: Kite is all here, Your Honor.
8	THE COURT: How much additional on direct do you
9	think you have?
10	MR. HEINRICH: I'd say a half an hour or less.
11	THE COURT: And then Mr. Weinberger, how much cross
12	do you think you I'm not trying to rush you, I just want
13	to get a feel for whether we can let the jury go a little bit
14	earlier today.
15	MR. WEINBERGER: I would say 45 minutes to an hour.
16	THE COURT: Okay. Thank you.
17	MR. DANE: We did the next witness is Dr. Schuetz
18	who does need to get on today.
19	THE COURT: Okay. So how long will how long is
20	his direct?
21	MR. DANE: Mr. Lawton is doing his direct. He's not
22	here. I think it should be on the order of 25 minutes,
23	something like that.
24	THE CLERK: Your Honor, are we ready?
25	THE COURT: Yes.

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1 THE CLERK: Would you all rise for the jury, please. 2 (Thereupon, the jury returned to the courtroom.) 3 THE COURT: Okay. We have our jury reassembled with counsel present and the parties. We are ready to proceed, 4 5 so... MR. HEINRICH: Good afternoon, ladies and gentlemen. 6 7 Welcome back. 8 BY MR. HEINRICH: O. Dr. Sullivan. 9 10 A. Thank you. Q. So let's recap. You were identifying at a high level the 11 12 two adjustments you make to account for differences between the MSK/Juno 2013 agreement and the Juno/Kite 2017 13 hypothetical negotiation. So why is the hypothetical 14 15 negotiation in October 2017? A. That is when Kite obtained FDA approval to begin selling 16 17 YESCARTA. Prior to that point in time, they were only 18 engaging in clinical trials. And clinical trials in this space does not give rise to infringement. So companies can 19 20 undertake as many clinical trials as they wish, and that 21 would not be considered infringement. Hence, the infringement does not begin until October 18th, 2017, upon 2.2 23 FDA approval. Q. All right. So you start with the MSK/Juno 2013 24 25 agreement. Let's talk about the terms of that agreement.

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Let's pull up Exhibit 924. All right. And let's go to page 1 2 What royalty rate is specified in this agreement? 3 A. The agreement specifies a running royalty rate of 7.25 percent payable on net sales. And this is the royalty 4 rate that is payable by the licensee, which is Juno, as well 5 as any sublicensee, for example, Kite, in terms of what would 6 7 be payable to MSK under this agreement as it's specified. 8 Q. So if Juno were to sublicense Kite at the hypothetical negotiation for 7.25 percent, how would that royalty be 9 10 divided between Juno and MSK? 11 A. All of that, all the 7.25 percent would go to MSK and 12 Juno would receive zero. 13 Q. All right. Did you also evaluate other payment obligations that Juno made upon entering the agreement that 14 15 are not tied to sales? 16 A. Yes. There are milestone success payments as well. 17 Q. All right. Let's discuss each of those. Can we pull up 18 924, page 15, 924.15? So can you walk us through these 19 milestone payments? 20 A. Yes. So this is section 4.6 of the agreement, and this 21 is setting forth development milestone payments. The first one is \$50,000 for completion of a phase I trial. 2.2 There is 23 also a payment of \$300,000 for a completion of a phase III trial, and a payment of \$3 million for receiving regulatory 24 approval. All three milestones which were achieved by Kite 25

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as of the time of the hypothetical negotiation. 1 2 Q. So let's maybe unpack that a little bit. At the time 3 Juno entered this agreement with MSK, did Juno pay these amounts immediately? 4 There was a payment obligation, yet the payments 5 were not made at that particular point in time, but rather 6 7 these get triggered across time. Q. And so why did you consider them in your analysis of the 8 upfront payment obligations for your license structure? 9 10 A. As of the time of the hypothetical negotiation, these 11 milestones would have already been triggered. You know, this 12 is -- you know, the same approach that Kite's expert Dr. Rao takes in terms of looking at which milestone success payments 13 have been triggered as of the time of the hypothetical 14 15 negotiation. Q. All right. So let's discuss the success payments that 16 17 you mentioned. Can we pull up PX241? Which is another 18 co-agreement between Juno and MSK. And if we turn to page 23, 241.23, what do we see here? 19 20 A. So this is part of the agreement between MSK and Juno. 21 And this is setting forth the success payments under that agreement. What it is showing is that the success payments 2.2 23 are based upon increases in equity value, such that the first payment, it's showing here as \$10, but as you can see, this 24 is in millions of dollars, so there is a payment of 25

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\$10 million when the equity value that the stock price increases by a multiple of 10. Then there is an additional \$70 million payment to get up to a total of \$80 million that the stock price increases 15-fold, and if the stock price increases 30-fold, then the entire payment is \$150 million under the success payment. Q. And why did you consider these success payments as part of your upfront in your royalty calculation? A. Well, for two reasons: One, they are a fundamental component of the agreement. This is, you know, a basis for why Sloan Kettering and Juno entered into this agreement. Secondly, all of these payments -- similar to the milestone payments I was mentioning a moment ago, all of these payments were also triggered. So when one takes a look at the increasing value of Kite's Series A stock, that increased well over 30-fold. And --MR. WEINBERGER: Objection, Your Honor. THE WITNESS: -- as a result, those would be triggered. THE COURT: Grounds? MR. WEINBERGER: Grounds on MIL. This is stock swap expressly. THE COURT: Overruled. BY MR. HEINRICH: Q. So in sum, how much of the upfront payment obligations

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not tied to sales did you include as your starting point for that component of the hypothetical license you proposed? A. \$153,350,000, so that consists of the milestone payments of 3,350,000, plus the milestone success payments that are listed here of \$150 million. Q. How does Kite's expert value these success payments in his analysis? A. Well, it's rather interesting, because while he takes a similar approach in looking at milestone payments, he uses a different agreement. Rather than using the Memorial Sloan Kettering/Juno agreement, he uses a different agreement that is not for the patented technology. Q. All right. Let's turn to the adjustments you make and getting to the details of those adjustments. If we can go to this slide, thank you, this is slide 27. How do you make your first adjustment? A. The first adjustment is based upon two agreements that were entered into by Juno; one of them with St. Jude, and the other one with Novartis. And by comparing those two agreements, it allows us to make an adjustment that gets us to early 2015 with Novartis as a party to the negotiation. There's still differences at that point, but it takes us to part of the adjustment. Q. Okay. So if --

MR. HEINRICH: So I would like to move into evidence

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1	Exhibit 401, the Juno/St. Jude agreement.
2	MR. WEINBERGER: No objection, Your Honor.
3	THE COURT: PX401 is received.
4	(Exhibit PX401 received.)
5	BY MR. HEINRICH:
6	Q. And is this Juno/St. Jude Children's Research Hospital
7	agreement one of the agreements you considered for your first
8	adjustment?
9	A. Yes. This is the agreement I was just referring to.
10	MR. HEINRICH: And then I'd move into evidence
11	PX928, the Juno/Novartis settlement sublicense.
12	THE COURT: Any objection?
13	MR. WEINBERGER: No objection, Your Honor.
14	THE COURT: Thank you. Received.
15	(Exhibit PX928 received.)
16	BY MR. HEINRICH:
17	Q. And is this the second agreement you considered for your
18	first adjustment?
19	A. Yes, it is. This is the other agreement I just referred
20	to.
21	Q. And if we can now turn back to the slides, why don't you
22	walk us through how you calculate this first adjustment?
23	A. So the agreement that Juno entered into with St. Jude in
24	2013 specifies a running royalty rate of 2.5 percent.
25	Subsequently, in 2015, Juno entered into a settlement
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agreement with Novartis that involved the very same patent, and the running royalty rate in that agreement is

4.75 percent. So you can see that what has occurred across time, going from 2013 to early 2015, as well as the difference in a fully -- thinking about entities that are doing research versus those that are developing a CAR-T product, there's a difference overall that goes from

2.5 percent up to 4.75 percent.

And that reflects a 90 percent increase relative to

And that reflects a 90 percent increase relative to the 2.5 percent. So that provides us with effectively what is the first adjustment.

Q. Now, the patent at issue in these agreements, is it the same Sadelain patent we're here to discuss?

A. No. It's a different patent. It does relate to CAR-T constructs, but it is a different patent. And it has different attributes, and different value in terms of licensing.

And you can see that straight away because the royalty rate in the St. Jude agreement is at 2.5 percent, where as the royalty rate in the Memorial Sloan Kettering/Juno agreement is higher at 7.25 percent.

So that tells us that there's a difference in the patents that are being licensed, but it is something that allows us to look at the relative ratio or the relationship.

O. All right. So you indicated that you did a second

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adjustment. And why did you feel the need to do a second adjustment? We're now at slide 29.

A. There are still fundamental differences between the agreement with Novartis in 2015 and the agreement that would result from the hypothetical negotiation in 2017 with Kite. And in particular, back in 2015, Novartis was still years away from being able to launch a product. They were in -- still in development phase. There was no overlap with the lead product candidates between Juno and Novartis. The agreement is also a settlement agreement. It settles litigation. And so it was resolving uncertainty, resolving litigation costs, especially in light of the impending intellectual public -- initial public offering, the IPO for Juno.

And there's also considerable evidence that Novartis had a questionable commitment across time to the CAR-T space. And for all of these reasons, that's very different from what's occurring at the hypothetical negotiation, which is in 2017, October of 2017, where we have Kite launching a product into the marketplace that's fully commercialized, and is the primary competitor of Juno.

- Q. So how does -- so does Dr. Rao, Kite's expert, do some of the same type of analysis?
- A. Well, here again, Dr. Rao and I agree that a starting point is the 2013 agreement with 7.25 percent as that rate.

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1 And we both make our first adjustment, I make a first 2 adjustment. Dr. Rao does that, too, although he does a 3 different approach to that adjustment so that he ultimately ends up at a royalty rate of 10.34 percent. However, he does 4 not account for any of the differences between Novartis in 5 2015 and Kite in 2017 as it relates to how it impacts Juno. 6 7 And that's what the second adjustment accounts for. 8 Q. So does Dr. Rao stop his adjustments at the Novartis/Juno 9 2015 settlement agreement? 10 A. Yes. That's right. 11 O. All right. So can you first explain at a high level your 12 adjustment number 2? A. Yes. So there's two real distinctions. In 2015, we have 13 Novartis versus at the hypothetical negotiation in 2017, we 14 15 have Kite. And that -- they have different competitive effects based upon what was being expected for competition in 16 17 the marketplace. 18 Second, the market had developed from 2015 to late So I account for both of those items by looking at the 19 relative competitive effects on Juno. 20 21 So I first look at what would have been the 2.2 competitive effect on Juno in 2015 from Novartis and then I 23 look at the competitive effect of Kite in 2017 on Juno. And I look at that relative difference. 24 Q. And so can you explain to us how you do that calculation? 25

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A. Yes. So first off, I look at the market share effects on Juno. These are based upon the financial models that were used by Juno for their business efforts, both for their product pipeline, all their business decision making, as well as for evaluating any sort of acquisition.

In those models, around the time of the Novartis 2015 agreement, Juno expected that Novartis would ultimately have a market share effect on Juno of 4.4 percent.

In contrast, fast forward to 2017 around the hypothetical negotiation, at that point in time, Juno projected through their financial models that Kite would have a market share effect of 12.5 percent, a greater market share effect on Juno.

- Q. How did you select the models that you used for this analysis?
- A. Based upon the use of those models by Juno in the ordinary course of business, looking at the completeness of the models, validating the work that was performed, as well as looking at which models were closest in time to each of these two events, the Novartis agreement in 2015, as well as models that are closest in time to the hypothetical negotiation in October 2017.

MR. HEINRICH: So I move into evidence Exhibits 1247 and 1248, which are summaries of the models.

MR. WEINBERGER: Your Honor, I don't believe they

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1 are summaries of the models, and they are not evidence. They 2 are just demonstratives that were prepared, I think, by 3 Dr. Sullivan. MR. HEINRICH: We offer them under section 1001 as 4 5 summaries of voluminous data. THE COURT: 1247 and 1248, are they in the exhibit 6 7 binders here? 8 MR. HEINRICH: Yes. I'm sorry. I meant to say section 1006, rather than 1001. 9 10 THE COURT: Lay some additional foundation. 11 BY MR. HEINRICH: 12 O. Can you explain what Exhibit 1247 is, Dr. Sullivan? A. Yes. So the financial models that I was referring to are 13 large and complex Excel workbooks. They have many different 14 tabs, many different tables. There is a lot that goes into 15 16 these. As you heard earlier, there can be several months' 17 worth of a team putting these together. And what I have done 18 is I have pulled out the key information to summarize those financial models in Exhibits 1247 and 1248. 19 MR. HEINRICH: With that, we move them into evidence 20 21 under Federal Rule of Evidence 1006. 2.2 THE COURT: Same objection? 23 MR. WEINBERGER: Yes, Your Honor, they are 24 selective, they are not complete. THE COURT: Let me just -- okay. These -- 1247 and 25

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1248 are being received into evidence. These are certain charts and summaries. Charts and summaries are only as good as the underlying evidence that supports them. You should therefore give them only such weight as you think the underlying evidence deserves. And with that, they are received.

(Exhibits 1247 and 1248 received.)

BY MR. HEINRICH:

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- Q. So how did you quantify the relative economic impact of Novartis on Juno in 2015 and Kite on Juno in 2017?
- A. So as you can see on the next slide, that I converted those market share effects into dollar effects, looking at it on a revenue basis. And I do that by looking at this on an indication basis, converting market share into number of patients, looking at the price per therapy at the various points in time, and performing an annual revenue impact estimate.

And that's what you'll see here on slide 33. So that in 2015, Novartis was projected to have an annual revenue impact of \$445 million on Juno over the course of the agreement. And in contrast, in 2017, Kite was projected to have an annual revenue impact of \$1.3 billion.

And so there's a relative difference, a ratio of -resulting in an increase of 192 percent. In other words,
the -- starting at 445 million, going up to 1.3 billion,

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that's a 192 percent increase, so that's the second 1 2 adjustment factor that I use. 3 Q. Can you help us understand, you know, whether this makes sense at a high level? 4 MR. WEINBERGER: Objection, Your Honor. 5 6 THE COURT: Grounds? 7 MR. WEINBERGER: Vaque. 8 THE COURT: Okay. Sustained. 9 BY MR. HEINRICH: 10 Q. So as a matter of just basic economics, would one expect 11 or not expect to have this difference between market impacts 12 in these two periods of time between these two competitors? A. Well, it makes sense as an economist, but I do think it's 13 common sense that back in 2015, Novartis was still years away 14 15 from a launch, and they were not the primary competitor. They were seeking to launch in a different indication, versus 16 17 Kite in 2017 launching right at the lead product indication 18 for Juno, and they're actually launching the product. So you would expect there to be a larger effect in 2017 from Kite 19 20 than you would from Juno -- from Novartis in 2015. 21 Q. Now, to be clear, are you doing a lost profits analysis? 2.2 A. No. This is not lost profits at all. This is just looking at the relative effect. And I use this relative 23 24 effect, and I apply that to royalties. So there is no capture here of revenue, there is no capture of profits. 25

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It's purely looking at the relative effects. And I use that 1 2 relative difference to adjust the royalties in the Sloan 3 Kettering agreement as the basis. Q. So let's talk about your calculations using these 4 adjustments. So let's start with your royalty rate. So can 5 you walk us through your calculation. And this is slide 34. 6 7 A. Yes. As noted, the 2013 agreement between Memorial Sloan Kettering and Juno specifies a running royalty rate of 8 7.25 percent. Adjustment number 1 is adding 6.5 percent to 9 10 that amount. 11 And how I get there is I take the 90 percent 12 adjustment factor from adjustment number 1, and I multiply that by 7.25 percent, and that yields 6.5 percent. So there 13 is an additional royalty for adjustment 1. 14 And then if we go to adjustment number 2, I take 15 16 that same base of 7.25 percent, I multiply it by the 17 adjustment factor of 192 percent, as I explained for 18 adjustment number 2, and that provides an incremental royalty rate of 13.9 percent. And so when I add these three 19 together, the 7.25, 6.5, and 13.9, that yields an ultimate 20 21 running royalty rate of 27.6 percent. 2.2 Q. And of that amount, how much -- so we imagine this 23 hypothetical sublicense between Juno and Kite. How much of 24 that royalty rate would go to MSK?

A. 7.25 percent would go to MSK. And the remainder, just

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over 20 percent, would go to Juno. 1 2 Q. So how does your proposed rate compare to Kite's 3 anticipated profit margins at the time of the hypothetical negotiation? 4 A. At the time, based upon the financial models prepared by 5 Kite for their Project Gold, that specifies profit margins 6 7 ranging from about 65 percent in the beginning going up to 8 over 80 percent profit margins by the time that the patent 9 would expire in 2024. 10 Q. So can we pull up Exhibit 156, and turn to page 25? Ιf 11 we can just highlight the gross profit margin up above. 12 And what is this document? 13 A. So this is the Project Gold report that was put together by Kite based upon their financial models that they had put 14 in place just a bit before the hypothetical negotiation. 15 And here, as I was noting earlier, the gross profit 16 17

And here, as I was noting earlier, the gross profit margins range in the beginning from about 66 percent and goes up to over 80 percent by the time that the patent would expire in 2024. And in dollar terms, the gross profits during this point -- during this time would be \$8.6 billion during the life of the patent up until it expires in August of 2024.

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Similarly, there is also a line here for operating profits that deducts other operating expenses, you know, selling expenses, executive salaries and the like. And the

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total during the life of the patent is 7.1 billion. So the 1 2 profits that were projected were just over \$7 billion on a 3 net operating basis. Q. All right. So let's take your royalty rate and come up 4 with the running royalty component of your damages opinion. 5 If we can go to our next slide. 6 7 So what -- what are you applying your royalty rate 8 to? A. The base -- the royalty base of net sales of YESCARTA 9 10 from inception in 2017 through the third quarter of this 11 year, the third quarter of 2019. Those sales are 12 \$604 million. And that amount of sales is not disputed in 13 the case. O. And are those worldwide sales? 14 A. Yes, they are. It includes global sales because all of 15 the production is performed here in the United States. So it 16 17 makes sense to include worldwide sales. 18 And applying the 27.6 percent rate to the sales of \$604 million provides for running royalties of 167 million. 19 Q. All right. So we saw that you have a second component of 20 21 your proposed royalty, which is an upfront component or a component not based on sales. Can you walk us through the 2.2 23 calculation there. A. Yes. The calculation is parallel to this or what I have 24

done for the running royalty rate. So as you'll recall, the

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1 milestone success payments total \$153.35 million. 2 results from the 3.35 million milestones and the \$150 million 3 success payments. So I apply the two adjustments to the 153.35 million 4 just the same. So I apply a 90 percent factor --5 MR. WEINBERGER: Excuse me, Your Honor. Before this 6 7 is published, I need to object for the record. 8 THE COURT: Go ahead. MR. WEINBERGER: This is contrary to the MIL ruling 9 10 regarding the relationship between the upfront payment and 11 the revenues and profits. 12 THE COURT: Thank you. 13 THE WITNESS: So I apply the same two factors to the \$153.35 million milestone success payments. I apply a 14 15 90 percent factor, and I apply a 192 percent factor. And when I do that math, just like I did for the running 16 royalties, the total amount is \$585 million for the upfront 17 18 component of the reasonable royalty. BY MR. HEINRICH: 19 Q. And what does that take into account? 20 21 A. So that is accounting for the financial harms and 2.2 benefits that are realized at the time of the hypothetical 23 negotiation. For example, right at the hypothetical negotiation, 24

by allowing Kite to launch with the license, by providing a

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license, providing them with permission to launch, that provides the first mover advantage for Kite. And it also provides the reciprocal harms to Juno at that time. And those become enabled or realized at the time of the hypothetical negotiation. Q. So you mentioned a few disagreements you have with Dr. Rao, Kite's damages expert. Can you summarize those for us here? While we both start for our running royalty with the 7.25 percent, his adjustments only go up to 10.34, because he's only making a partial adjustment, taking us to Novartis in 2015. And while I disagree with how he does his adjustment, that part aside, he has not accounted for any of the other differences between Novartis in 2015 and Kite in 2017. He also ignores the bargaining position of the parties in 2017. So at the hypothetical negotiation, it would be recognized that Kite no longer had options. They had tried to license the patent and did not have it. They had tried alternatives and design-arounds, and that did not

Q. And what are your total proposed damages in this case,

of Juno relative to Kite.

work. And thus, at the point in time of the hypothetical

negotiation, there would be a very strong bargaining position

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1	sir?
2	A. So adding together the running royalty component of
3	\$167 million to the upfront component of the reasonable
4	royalty, which is \$585 million, those together provide a
5	total reasonable royalty of \$752 million.
6	Q. And why do you think that that is reasonable in this
7	case?
8	A. Well, for a number of reasons. But the technology
9	underlying the Sadelain '190 patent is fundamental to
10	YESCARTA, and at the hypothetical negotiation, that license
11	enables Kite to launch into the marketplace and obtain the
12	benefits financially of doing so.
13	MR. HEINRICH: Thank you very much, Dr. Sullivan.
14	THE COURT: Thank you.
15	Mr. Weinberger?
16	MR. WEINBERGER: If I can just have a moment to set
17	up, please.
18	CROSS-EXAMINATION
19	BY MR. WEINBERGER:
20	Q. Good afternoon, Dr. Sullivan.
21	A. Good afternoon.
22	Q. You testified that the starting point for your analysis
23	was the license agreement between Memorial Sloan Kettering
24	and Juno in 2013, correct?
25	A. That is the starting agreement. My analysis really

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starts before then by looking at the marketplace, the role of the technology, importance of the patents -- of the patent, things of that nature. But then once I turn to the market approach, the starting point within that is indeed that agreement. Q. In fact, you concluded that the only agreement in this case that could be appropriate to use as a comparable and then adjust was the Memorial Sloan Kettering agreement, correct? A. That is right. The other two agreements can be used as relative amounts, but I do not think they would be appropriate as a starting point. Q. There are no other comparable licenses in this case according to you, correct? A. Not ones that could be reasonably and reliable adjusted to account for the differences between those agreements and the hypothetical negotiation. Q. Now, you have testified that there is a set of two other agreements involving related technology that you used to establish and quantify certain adjustments, correct? A. That is right, the St. Jude agreement and the Novartis settlement. Q. Right. And the St. Jude agreement -- we can put up demonstrative number 1, which I have given to counsel. And these are just your calculations, so there is nothing in here FILED: NEW YORK COUNTY CLERK 03/24/2023 11:45 PM INDEX NO. 451625/2020 NYSCEF DOC. NO. 2:175CV-07639-GW-KS Document 605 Filed 12/18/19 Page 105 of 228 Page 1D NYSCEF 103/24/2023 #:26186

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1	that you haven't said already.
2	The St. Jude license has a royalty of 2.5 percent,
3	correct?
4	A. That's right.
5	Q. And the Novartis license has a royalty rate of
6	4.75 percent, correct?
7	A. That's right.
8	Q. And as we discussed, the MSK/Juno agreement has a royalty
9	of 7.25 percent, correct?
10	A. Yes, that's right.
11	Q. And your royalty is 27.6 percent, correct?
12	A. Yes, correct.
13	Q. So that's almost four times higher than the MSKCC Juno
14	license, right?
15	A. Almost, yes.
16	Q. And it's about six times higher than the Novartis
17	license, correct?
18	A. Um, about that, yes.
19	Q. And it's about 11 times higher than the St. Jude license,
20	correct?
21	A. Yes, that's right.
22	Q. And there are no other agreements that you have opined
23	about in your report that are comparable in this field of
24	technology, correct?
25	A. No, not in the way that I have defined and used the term
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comparable, that's right. 1 2 Q. All right. So you made two adjustments that you've 3 testified about, and I'm just going to put up another chart summarizing the numbers on that. And again, this is not 4 controversial. It's the same numbers that you have given the 5 6 jury. 7 Could we post number 2? 8 So first you did what you called a CAR-T developer adjustment, and that got you from 7.25 to 13.78 percent, 9 10 correct? 11 A. Yes, that's right. 12 O. So you close to doubled the royalty rate in the MSKCC agreement with your first adjustment, right? 13 A. It's a 90 percent increase, that's the 90 percent factor. 14 15 Q. And then you made a second adjustment which you called a Kite competition adjustment, correct? 16 17 A. Yes, that's right. So that reflects the second 18 adjustment, the 192 percent factor. Q. And then that got you to 27.6 percent. I actually think 19 20 these numbers are reversed. It says 26.7, but I think it's 21 actually 27.6, correct? A. The answer is 27.6. 2.2 23 Q. Right. So you first -- so you basically took the highest rate of the three licenses you looked at, and you basically 24

quadrupled it, correct?

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1	A. No, that is not quite right. While in the end the
2	numbers are that, but I also examined a great number of
3	different license agreements, so I considered a large number,
4	but of these three, this has the largest royalty rate.
5	Q. I'm just asking about the math. The math is that you
6	almost quadrupled the highest license in the three that you
7	examined closely, correct?
8	A. The exact factor is 3.82, so it's not quite four, but
9	close enough.
10	Q. All right. Now, you also testified to an upfront
11	payment, correct?
12	A. Yes, that's right.
13	Q. And that was based upon the payments called for in the
14	MSKCC general license, correct?
15	A. Um, the agreements between Juno and Memorial Sloan
16	Kettering.
17	Q. Now, to be clear, you testified to two different types of
18	payments. The first one were milestone payments that totaled
19	\$3.5 million, correct?
20	A. 3.35 million.
21	Q. 3.35. Thank you.
22	And the second one was what you called success
23	payments, correct?
24	A. Yes. That is how they are termed in the agreement.
25	Q. And the success payments in the agreement basically

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provide that if Juno stock got to certain levels, 10 times higher, 20 times higher, 50 times higher, then payments would be made to Memorial Sloan Kettering, correct? A. Um, it's 10 times, 15 times, and 30 times. Under that agreement, Juno is the licensee, whereas at the hypothetical negotiation Kite is the licensee. Q. Understood. But what I want to just make sure is clear is that the success payments in the Memorial Sloan Kettering agreement were based upon the price of Juno stock, not Kite stock, correct? A. That's right, you know, similar to the running royalty would be applied to Juno sales not Kite sales --Q. Could you please just answer my question, sir. You know how this is done. You are a professional witness. have a short time period. We are trying to get the jury out of here. Just answer the question. THE COURT: That's argumentative. BY MR. WEINBERGER: Q. All right. So Juno's stock did not reach the levels by October 2017, which would have triggered the \$150 million payment you testified about, correct? A. Yeah, I'd have to think about exactly when those were triggered. It probably didn't occur until early 2018. Q. And in fact, by 2017, in October, the date of the hypothetical negotiation, the only payment that would have

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1 been due to MSKCC under that stock provision was \$10 million, 2 correct? 3 A. I don't recall the exact timing, that may be right, but I actually was thinking there had been further triggers at that 4 point. But I don't have that -- that level of precision in 5 6 my memory. 7 Q. So you can't dispute right now that the Juno stock had 8 reached the level where only \$10 million was due to MSKCC, not 150, correct? 9 10 A. That's right. 11 Q. So the way you got to 150 was that you substituted Kite's 12 stock price for Juno's stock price, correct? 13 A. No, not at all. Q. Well, how did you get to \$150 million based upon a Juno 14 15 stock success factor that was not reached? 16 A. You are confusing Juno and Kite. So at the hypothetical 17 negotiation, we specify what would be the outcome of that 18 hypothetical negotiation. Starting with an agreement between Sloan Kettering 19 20 as the licensor, Juno as the licensee. But it's different at 21 the hypothetical negotiation, because there, Juno is the licensor and Kite is the licensee. 2.2 23 Q. All right. So but the question is, you calculated 24 \$150 million based upon the Kite stock price in October 2017, 25 correct?

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1	A. It is based upon the triggers on the success payments.
2	Just like what was done by Kite's expert, Dr. Rao.
3	Q. Yes, but the trigger was the Kite stock price that you
4	used, correct, not the Juno stock price?
5	A. Applying the agreement and applying the terms, it applies
6	to Kite, it applies to Kite sales, and it applies to the
7	performance of YESCARTA.
8	Q. So the answer to my question is yes, you used the Kite
9	stock price to determine whether the success payments were
10	triggered, correct?
11	A. I would put it differently.
12	Q. Well, that's what you presented in your slide this
13	morning as to how the \$150 million was triggered. Right?
14	A. I put it differently.
15	Q. The Juno stock price I just want to make sure we're
16	clear on this. The Juno stock price never got to the levels
17	that would have triggered that \$150 million payment to
18	Memorial Sloan Kettering, correct?
19	A. Um, in effect, yes, given the acquisition.
20	Q. October 2017 the Juno stock price was nowhere near the
21	level that would have triggered \$150 million payment to
22	Memorial Sloan Kettering; yes or no?
23	A. I would have to disagree with you.
24	Q. You think the Juno stock price did reach a level that
25	triggered \$150 million success payment?
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A. Ultimately, as I understand it, there were payments and there were monies received by Sloan Kettering of approximately \$150 million. Q. Excuse me, but I'm asking as of October 2017, and I think we have been through this already. A. It --Q. Let me finish the question. As of October 2017, you agree that the Juno stock price was nowhere close to the multiple that would trigger \$150 million success payment, correct? A. Well, in all fairness, some of your questions are October 2017, others aren't. O. If I'm not clear, let me know. I want to be clear. A. Yeah. And I just can't agree with this characterization of nowhere near. I don't have the precision of time of when it hit, but it was getting darn close around that time, and I just don't recall that. So I can't say nowhere near. Q. So if hypothetically that payment that would have been earned at that time was only about \$10 million and not \$150 million, you would agree that would be nowhere close, right? A. There is a significant difference between 10 and 150. Q. Okay. Now, so if that was true, then the real world payment to Memorial Sloan Kettering is 10 million -- strike that question. Let's move on.

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1 Now, Juno contends that both companies were using 2 the '190 patent in their product development, correct? 3 A. Both Juno and Kite. Q. And both companies -- Juno used the patent in the JCAR15 4 product that it tried to bring to market, correct? 5 That's right. 6 A. Yes. 7 Q. And there were patient deaths that the jury has heard 8 about, and the clinical trials were halted by the FDA, 9 correct? A. There has been testimony along those lines. 10 11 Q. Kite figured out how to make a product that was approved 12 by the FDA as safe and effective, correct? 13 A. Well, Kite did launch a product. I can't speak to exactly the development process, given the relationship with 14 15 NCI. Q. The question was it figured out how to make a product 16 17 that was approved by the FDA as safe and effective; is that 18 right? A. It has made a product that is approved by FDA. 19 Q. So Kite's stock price would reflect the success that they 20 achieved that Juno never achieved, correct? 21 2.2 A. Close. It reflects the successes of Kite, it does not --23 and YESCARTA, but not with the -- what you were saying with 24 relative to Juno.

O. Now, let me just ask you a few questions about the last

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1	item you talked about, which is the profit. And first, it's
2	correct that the numbers you've given the jury, the upfront
3	payment on the royalty rate are only through the time of
4	trial, correct?
5	A. The upfront payment is a single payment. It doesn't get
6	made multiple times. So that applies throughout the term.
7	Q. That's not what you said in your report, is it, sir?
8	A. Um, yes, I said it's a single, one-time upfront payment.
9	The definition of an upfront payment is that it's paid up
10	front and paid once. It's not a multiple or recurring
11	payment.
12	Q. But you understand that Juno has reserved the right to
13	come back and ask Kite for
14	MR. HEINRICH: Objection.
15	THE COURT: Let him let him finish his question.
16	And the question of counsel
17	MR. WEINBERGER: I'll withdraw it, Your Honor. I'll
18	ask another question.
19	THE COURT: All right. Thank you.
20	BY MR. WEINBERGER:
21	Q. Would you look to paragraph 54 of your report, which
22	should be in your binders there.
23	A. Sure. Give me just half a moment.
24	Q. Sure.
25	A. Yes, I'm there.

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1	Q. All right. Now you're going to have to wait for me.
2	Sorry.
3	THE COURT: What number is it?
4	MR. WEINBERGER: Paragraph 54.
5	THE WITNESS: Page 31.
6	BY MR. WEINBERGER:
7	Q. And in this paragraph 53, you state, "The reasonable
8	royalty discussed herein," which includes both the upfront
9	payment and the running royalty, correct?
10	A. Yes. So this is referring to
11	Q. Just that statement, "reasonable royalty," refers to
12	just the to both the upfront payment and the running
13	royalty, correct?
14	A. Ultimately, reasonable royalties do include both
15	components.
16	Q. And in your report, you stated, "The reasonable royalty
17	discussed herein provides a reasonable estimate of damages to
18	Juno caused by Kite's infringement for the period from 2017
19	to Q4 to 2019 Q1," and you updated that, sir, to this last
20	quarter, "Royalties discussed herein do not represent
21	compensation for any damages or harm beyond 2019 due to the
22	infringement by Kite." Is that correct?
23	A. Yeah. That's right. So the one piece here that you
24	omitted is that's for the period for which Kite has provided
25	sales data.

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Q. Well, we can't provide sales data for 2020, it hasn't 1 2 happened yet. 3 A. Exactly. Q. Okay. But you also say in paragraph 54, "Royalties 4 herein only capture damages through 2019 Q1, and as such, 5 represent a period of time in which Juno does not have a 6 7 commercialized product in the market," correct? 8 A. That's true. Q. And then you say at the bottom of that paragraph, 9 10 "Damages for the period after JCAR17's launch may include 11 lost profits on JCAR17 sales due to Kite's infringement, in 12 addition to a reasonable royalty," correct? A. Yes. That's right. And as you will recall from earlier, 13 the statute provides for damages no less than a reasonable 14 royalty. And the reasonable royalty is far less than what 15 lost profits would be. So there's a potential, as a legal 16 17 matter, that subsequently perhaps there could be a situation 18 where lost profits could be in addition to reasonable 19 royalties. In other words, there could be a difference 20 between the two, and the delta may be available later on if 21 that's ever determined in a separate court action. 2.2 Q. So the point I'm making is although you've talked a lot 23 in your testimony about the products that Juno wants to bring to market, JCAR17, and the nature of competition that would 24 happen at that time, that is not -- the lost profits on such 25

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1 products are not included in the numbers you put before the 2 jury, correct? 3 A. There is a delta between lost profits and reasonable royalties. The royalties capture competitive effects, but 4 isolated to the value contribution of the technology separate 5 and apart from other items, lost profits is more expansive. 6 7 It's not apportioning to the contribution of the technology, 8 it's looking at the whole kit and caboodle, it's looking at the entire profits across the entire product. 9 10 MR. WEINBERGER: Your Honor, I move to strike 11 everything in the answer after yes. 12 THE COURT: I don't think he said yes. 13 MR. WEINBERGER: Well, then I'll ask it again. 14 BY MR. WEINBERGER: 15 Q. The royalty damages that you've put before the jury do not include any damages that Juno might seek for lost profits 16 17 after it comes to the market with a competitive product, 18 correct? A. For the reasons I just described, there could be a delta 19 20 between --21 Q. Sir, please, is it correct or not? 2.2 A. For the reasons I described to you. 23 Q. So the answer is yes? 24 A. For the reasons I described, yes. 25 Q. So that means if you're Kite and Gilead sitting at the

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hypothetical negotiation, trying to determine what you might be willing to pay for this license, you have no idea what your exposure might be after 2019 if Juno comes back and says now we are going to sue you for lost profits, correct? A. No. That is fundamentally wrong. And the reason is that at the hypothetical negotiation, if there's a license, that means there is permission. That would mean that Kite would be granted permission to use the patent, and thus there would not be a future legal proceeding. Q. Oh, I understand that they have permission to use the patent, but Juno can come back and claim that they need to pay a lot more money for the license, correct? A. No. Again, you are fundamentally wrong, because you are confusing an outcome of a hypothetical negotiation versus a court action that might result in damages in the face of Kite's actual infringement. Q. So just to clarify, this upfront payment and the royalty rate you've testified to, is that going to be what Kite will have to pay until the end of the patent term, which is 2024? A. I have not made a determination of post-judgment royalties, and there is a distinction, as I understand it, in the law. I'm an economist. But my understanding is the law is that once there is a finding of infringement, and if that infringement is willful, there can be other forms of royalties that might be available. And I have not been asked

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to do that. I have been looking at what would be the royalty rate resulting from the hypothetical negotiation. So again, it's apples and oranges between a court action versus a hypothetical negotiation that's not a court action. Q. All right. So the bottom line is that Kite would not know in 2017 at the hypothetical negotiation how much more it might have to pay in lost profits after Juno comes to market with a product, right? A. That is incorrect. At the time of the hypothetical negotiation, there would be certainty and resolution in the hypothetical. Q. But Juno could still claim lost profits damages when their product comes to market, right? A. Not in the hypothetical world, but in the actual world, in the face of the actions and activities undertaken by Kite. Q. All right. So let me move on to another subject. Let's talk about a basic principle, patent damages. You believe that a reasonable royalty should separate out the value contribution of the patented technology from other contributions, correct? A. Yes. I agree. Q. And you've written articles that say that, correct? A. I'm sure that I have. Q. And you've said in those articles that the apportionment

between the patentees -- of the patentee's damages between

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1 the patented feature and the unpatented features must be reliable and tangible, not conjectural or speculative, 2 3 correct? A. I don't recall those exact words, yet I do agree with 4 5 them. Q. Okay. Now, you showed the jury a slide from 6 7 Ms. Tomasello who talked about the importance of the CAR 8 construct? A. Yes, that's right. 9 10 Q. I think she said something like there would be no product 11 without the construct, correct? 12 A. That's right. 13 Q. And you relied on that in part for your opinion about the value of the patent, right? 14 A. It's one element that goes towards why the patented 15 technology is fundamental. 16 17 Q. Now, she also said that there were other elements that 18 were important for this product, correct? There are other factors, things like manufacturing 19 and lymphodepletion and administration. 20 21 Q. And if you don't have clinical protocol and results for a 2.2 product like this, you don't have a product, do you? 23 A. Um, not a commercialized product. Q. And if you don't have good manufacturing processes that 24 can safely and effectively manufacture the product, you don't 25

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have a product, correct? 1 2 A. You're not going to have a successful product. 3 Q. Right. And if you don't have FDA approval, you're certainly not going to have a product, correct? 4 A. Not a commercialized product. 5 Q. And the Juno product didn't have any of those things, did 6 7 it? I'm talking about JCAR15 right now, the product that 8 utilizes the '190 patent construct. Is that right? A. Um, I think I'm going to have to disagree with you on 9 10 that. 11 Q. Well, you're not disputing that the FDA that Juno's stock development of JCAR15 in 2017, right? We kind of talked 12 about that already. 13 A. They did cease continuing trials of JCAR15. 14 15 Q. And is it true that Juno determined that it would have to make protocol modifications and process improvements if it 16 17 wanted to pursue JCAR15 further, right? 18 A. Um, I believe those were items that Juno was considering in its next course of action. 19 Q. Well, in your report, you state, "Although Juno believed 20 21 that it could have proceeded with JCAR15 in clinical testing with certain protocol modifications and process improvements, 2.2 23 Juno would have first needed to conduct a phase I clinical trial to establish preliminary safety and to select a 24 recommended phase II dose with those improvements;" is that 25

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1 right? 2 A. That sounds right. 3 Q. And protocol modifications means changes to the clinical protocol, correct? 4 5 A. Yes. Q. And process improvements means improvements to the 6 7 manufacturing process, right? 8 A. It's a broader term. There's a lot of aspects to the overall process. It's not just manufacturing. So there's 9 10 different elements that could have been altered. 11 Q. And you are not -- it's not your opinion that Kite did 12 not make contributions to the YESCARTA product, beyond the 13 patent, is it? A. It is not my opinion that they did not make 14 15 contributions. Q. And you agree that the construct under the patent is not 16 17 sufficient to get an approved product, correct? 18 A. Not in and of itself. It's a fundamental basis, but it's 19 not sufficient. Q. And we know it's not sufficient because Juno using it was 20 21 unable to complete the work necessary to get an approved 2.2 product, right? A. Um, not quite right. They did not continue to proceed 23 24 down that path, but as we just noted a moment ago, that doesn't mean they were unable; rather, they chose. 25

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Q. And among the factors that go into providing a 1 2 commercialized product are the ones we talked about; 3 manufacturing process, clinical protocols. So far is that correct? 4 A. There is a variety of things that go in; manufacturing, 5 administration, lymphodepletion regimens. There is even a 6 logistical platform called Kite Connect that helps with 7 8 scheduling of patients. I mean, all of these things, you know, go towards the sales of the product, but they are not 9 10 the demand drivers. 11 Q. When you say they are not the demand driver, you don't 12 think that a doctor knowing that a product has been safely manufactured, that the genetic engineering has been done 13 correctly, that the FDA has approved the product, you don't 14 think that drives demand for doctors to prescribe it for 15 16 their patients? 17 A. While I think some of those items are factors, I don't 18 think they are the drivers of demand. Now, you did throw in there the FDA approval, and certainly FDA approval is 19 20 important, but that is also based upon the efficacy of the 21 product, which as I understand it is based upon the Sadelain CAR construct. 2.2 23 Q. Well, the FDA approval encompasses all the things that I have been talking about, doesn't it; the manufacturing the 24 lymphodepletion, the clinical protocols, all of it, correct? 25

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1 A. All of that would be submitted to FDA. 2 Q. And it's not your opinion that doctors care which 3 construct is used in the drug, do they? A. It is kind of funny, because it's not the name of the 4 construct itself, it's whether the construct works. And 5 here, the basis for why Kite's YESCARTA works is because of 6 7 the Sadelain CAR construct. 8 Q. Even though Juno couldn't make it work, correct? A. Again, I'm going to have to disagree with you on that for 9 10 the reasons I described. 11 Q. That's what this is all about, disagreements. 12 So I want to turn now to your calculation of the 13 first adjustment you made, the CAR-T developer adjustment. Can we put back the first demonstrative. 14 15 And, again, just to remind the -- is that up? All 16 right. 17 And just to remind the jury, you multiplied the 18 royalty rate in here by almost double to adjust the MSK license to make it look more like a license between two CAR-T 19 developers, right? 20 21 A. Um, I applied a 90 percent increase, which is a factor of 1.9, close to a factor of 2, to get to a point of the effect 2.2 23 of a CAR-T developer of Novartis in 2015. Q. And you applied the CAR-T developer adjustment because 24

Memorial Sloan Kettering is not a CAR-T developer, right?

25

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1	A. That is part of the reason.
2	Q. But Novartis is a CAR-T developer, correct?
3	A. Yes, they were.
4	Q. And the Novartis license that was entered into, I think
5	in 2015, is closer in time to the hypothetical license than
6	the MSKCC agreement, right?
7	A. Yes, exactly.
8	Q. So you could have just looked at the Novartis license,
9	which already has a competitor as the licensee, and adjusted
10	that, right?
11	A. Well, one could do that, and that would be inappropriate.
12	Q. And one of the reasons you say it's inappropriate is
13	because it was a settlement; is that correct?
14	A. There is multiple reasons. It's a different technology.
15	You can see that from the underlying rates with the St. Jude
16	Research Hospital of 2.5 percent versus 7.25. It's also a
17	settlement of litigation. And there are other reasons.
18	Q. All right. Let's talk about the difference in
19	technology. What was licensed by St. Jude to Juno and then
20	from Juno to Novartis was what's called a 4-1BB construct,
21	correct?
22	A. It's a patent that relates to a 4-1BB construct.
23	Q. And that is the construct that Novartis used in
24	developing its product, correct?
25	A. It is, as I understand it, the construct for KYMRIAH,

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1	which is the Novartis product.
2	Q. And that is also the construct that Juno is currently
3	using to try and get JCAR17 approved, correct?
4	A. Yes, that is right. Let's be careful not to confuse in
5	this instance a patent with a construct, because the
6	underlying patent is different than the '190 Sadelain patent.
7	Q. But you understand that without a license Novartis could
8	not have used the 4-1BB construct, right?
9	A. I would think that's right.
10	Q. So
11	A. I
12	Q. That's fine.
13	A. No, I just need to qualify as I think it through. I
14	would have to think about that.
15	Q. You don't know?
16	A. Not right this moment.
17	Q. But you really can't testify here as comparing the 4-1BB
18	to the '190 patent which one is more valuable, can you?
19	A. So there is two distinctions there. Again, you are
20	confusing a construct with a patent. The '190 patent is what
21	provides the CAR construct. 4-1BB is a different construct.
22	It's not a patent.
23	Q. Now, let me ask you about this settlement issue that you
24	raised. You're you've written articles about cases from
25	the federal circuit numerous times, right?

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1	A. Multiple times anyway.
2	Q. And the federal circuit is the court of appeals that
3	specializes in patent cases, correct?
4	A. That's right.
5	Q. And in your report, you quote from a case called ResQNet
6	vs. Lansa. It's footnote 481 of your report.
7	A. I'm familiar with the case.
8	Q. And it's correct in that case that the Court found the
9	most reliable license in the record arose out of a litigation
10	settlement, correct?
11	A. In that case, yes, that's right.
12	Q. They said there is no rule that you can't use a
13	litigation settlement agreement and adjust it to come up with
14	a comparable royalty, is there?
15	A. I'm not aware of a blanket rule in that regard.
16	Q. Okay. Let's move to the second adjustment you made. And
17	this second adjustment basically again doubled the amount of
18	the royalty from 13.78 to 27.06, correct?
19	A. It's a 192 percent increase, so a factor of 1.92.
20	Q. So starting from the beginning, we are now about four
21	times higher than the MSKCC license, right?
22	A. The factor overall factor is 3.82, so just under 4.
23	Q. Right. And that adjustment is to account for competition
24	between Kite and Juno which you claim is more intense, more
25	important than the competition between Novartis and Juno,

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1 right? 2 A. It's the difference between the competitive effect of 3 Juno -- of Novartis in 2015, early 2015, relative to the competitive effect of Kite in late 2017. 4 Q. But talking about competitive effects, Juno and Kite are 5 not actually competitors in the market for sale of CAR-T 6 7 products, correct? 8 A. They are competitors on multiple fronts. O. My question was they are not currently competitors in the 9 10 market currently for the sale of these products to doctors, 11 correct? 12 A. Maybe I misheard you. Are you referring to Kite? 13 Q. Kite and Juno. A. Kite and Juno. 14 15 Q. Correct. 16 A. Okay. 17 Q. Juno doesn't have a product. 18 A. They have a product. It's not for sale in the marketplace. It's being used in clinical trials. So yes, 19 20 there is a product out there. 21 Q. Fair enough. A. And yes, they do compete. They compete vigorously. They 2.2 23 are competing for patients. They are competing for clinical trial sites. They are competing for investment. They are 24 competing on very -- on a myriad of dimensions. 25

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1	Q. The one thing they are not competing for, sir, is sales,
2	correct?
3	A. Juno is not selling, so they are not competing in terms
4	of dollars. They are competing for the clinical trials. So,
5	in other words, they are directly competing for providing
6	that therapy to a patient.
7	Q. You agree that they are not competing for sales, right?
8	A. Not for dollar sales.
9	Q. And there is no guarantee, sir, that JCAR17 will ever
10	come to the market, correct?
11	A. Not a guarantee. I mean, the marketplace widely views it
12	as very, very likely.
13	Q. But there is no guarantee, correct?
14	A. Correct.
15	Q. So Novartis has been out there for the last year and a
16	half and is out there right now paying Juno less than
17	5 percent royalty for selling KYMRIAH to treat the DLBCL
18	indication, correct?
19	A. There are some sales of KYMRIAH for DLBCL as I recall.
20	Q. But in your opinion, Kite should have been paying Juno
21	during the same period approximately 27.6 percent for selling
22	YESCARTA to treat the same indication, correct?
23	A. Yes. It's a different patent at a different point in
24	time.
25	Q. All right. Let's talk about the nature of competition

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from Novartis in 2015. 1 2 One of the reasons you doubled -- almost doubled the 3 royalty rate from the first adjustment is because you believe that Novartis was not a formidable competitor in 2015, right? 4 A. It's the relative competitive effects. So I -- you know, 5 the analysis compares the relative effect of Novartis in 2015 6 7 to the effect of Kite in 2017. 8 O. Okay. So, in fact, in your slide we can put up -- it's 29, I think. 9 10 You said -- among the things you said to distinguish 11 Novartis from -- in 2015 from Kite in 2017 was questionable 12 commitment to CAR-T, correct? 13 A. That's right. Q. All right. Now, at this point in time Juno was a company 14 15 that was -- had a rough value of about \$800 million, right? A. I'll take your word for it on that. 16 17 Q. It's in your report. 18 A. Even better. Q. Okay. And you are aware that Novartis is one of the 19 20 biggest pharmaceutical companies in the world, right? 21 A. I am. O. It's more than 200 times bigger than Juno, correct? 2.2 23 A. At that time? 24 Q. Yeah. A. Um, that's probably about right. I think Novartis was 25

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1	over 200 billion.
2	Q. Right. And your opinion is that Juno wasn't too worried
3	about competition from Novartis, right?
4	A. Um, no. There are competitive effects. In fact, if you
5	recall adjustment 2, it's comparing an annual effect of 445
6	million to 1.3 billion. So I'm not saying Novartis had a
7	zero effect. But I'm quantifying the distinction or the
8	difference between Novartis in 2015 and Kite in 2017.
9	Q. What you are saying you said that Novartis had a
10	questionable commitment to CAR-T, right?
11	A. Yes, that's right.
12	Q. And you testified that Juno was not as worried about
13	Novartis as they were about Kite in 2015, right?
14	A. Juno is not as worried about Novartis in 2015 as they
15	were about Kite in 2017.
16	Q. All right. So could you turn in your binder to
17	Exhibit 253, please.
18	MR. HEINRICH: I need a binder.
19	MR. WEINBERGER: I'm sorry.
20	THE COURT: This is DX253?
21	MR. WEINBERGER: Yes, Your Honor.
22	THE COURT: Thank you.
23	MR. WEINBERGER: If this isn't in evidence, I'm not
24	sure, frankly, I move to admit it.
25	THE WITNESS: Would you all mind providing me with a

1	copy?
2	MR. WEINBERGER: Do you have
3	THE WITNESS: I do not have a binder.
4	MR. WEINBERGER: It should be in one of your binders
5	there, an exhibit binder, 243.
6	THE WITNESS: No.
7	MR. HEINRICH: I don't believe he has defense
8	exhibits.
9	MR. WEINBERGER: My apologies.
10	THE COURT: Any objections to 253 being received?
11	MR. HEINRICH: No objections.
12	THE COURT: 253 is received.
13	(Exhibit DX253 was received.)
14	BY MR. WEINBERGER:
15	Q. And the cover of this document is an e-mail we can
16	publish that e-mail from Hans Bishop, who was the CEO of
17	Juno and who testified here this morning, in November 2014,
18	right?
19	A. That's what it looks like.
20	Q. And this was a presentation that was made to the CEO of
21	the company, correct?
22	A. I do not know.
23	Q. Now, over on the page that is the second page, which
24	bears the number 0002, it states, "Juno is in a very
25	competitive environment. Novartis is formidable, two-year

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head start and is committed to this space." Correct? 1 2 A. Yes. Let me just get my bearings here real quick. So 3 this was -- okay. So this was back in 2014. Yes, I see 4 that. Q. And that accurately reflects Juno's mindset in 2015 right 5 around the time that it granted Novartis a license for the 6 7 patents that we talked about earlier, right? 8 A. Well, I can't speak to the mindset of Juno from this document. I don't recall it. Yet it's not inconsistent with 9 10 how I have approached my analysis. 11 Q. So you agree that Juno believed that Novartis was a 12 formidable competitor with a two-year head start? A. It's not inconsistent with my view and my analysis. 13 Again, just going back to the point that I'm looking --14 15 Q. Could you just answer the question, please. 16 A. Fair enough. 17 Q. Appreciate it. 18 A. Fair enough. 19 Q. We would all appreciate it. 20 So you also said in your slide, talking about 21 Novartis in 2015, that there was no overlap with Juno's lead 2.2 profit candidates, right? 23 A. That's right. 24 Q. And by that you mean Juno thought that Novartis was focused on pediatric leukemia which was not Juno's focus, 25

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1 correct? 2 A. No. It was a simpler point that Novartis wasn't leading 3 with 3L DLBCL. Q. That's what I meant. If I didn't state it correctly, 4 your point was that Novartis was focused on pediatric 5 leukemia and Juno was focused on DLBCL indication, right? 6 7 A. No. You're -- it may be a subtle difference for you, but 8 focusing versus what a lead product candidate can be a little bit different. I was just taking the simpler point of the 9 lead product candidate. 10 11 Q. Let me ask you this: Juno was projecting in this time 12 period that Novartis would eventually enter into the DLBCL 13 space, right? 14 A. Yes. Q. And, in fact, the license wasn't limited to the one 15 indication; they had the freedom to develop any indication 16 17 they pleased, right? 18 A. I do not recall any restrictions in the license agreement for particular indications. 19 Q. I'm just trying to consolidate, so just give me one 20 21 second. And in 2015, isn't it correct that Juno did not have 2.2 23 a belief that it was going to beat Novartis to market with its JCAR17 product in third-line DLBCL? 24 A. I'm not sure I fully heard that right, yet -- my 25

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1 understanding is that Juno was not expecting to beat Novartis 2 to market for DLBCL. 3 Q. Right. And you never even -- you never looked for any evidence that suggested that at any time since the Novartis 4 license Juno thought it was going to beat Novartis to market 5 in third-line DLBCL, right? 6 7 A. Well, it depends upon the point in time. So if we are 8 focused on the hypothetical negotiation, that's, you know, a different point in time. 9 10 Q. Well, I'm talking about 2015. From 2015 forward, there 11 wasn't any evidence that suggested that Juno thought it was 12 going to beat Novartis to market in third-line DLBCL, right? 13 A. Sorry. Maybe I misheard you earlier. I thought you were focused on the hypothetical negotiation. 14 15 Q. No, starting in 2015. A. Back in 2015, um, there were different views on exactly 16 17 how that would proceed. 18 Q. And you've seen Juno projections showing that they expected to be approved in an indication well after Novartis, 19 20 right? 21 A. There are different projections that go different ways. 2.2 Depends again on the point in time and whether one is just 23 looking at company-announced information versus private 24 estimates. Q. And I was actually going to come to that next, because 25

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when you are dealing with a company like this which has people doing financial projections and modeling, there are a lot of different projections that say a lot of different things, right? A. There are different projections at different points in time. Q. And the basis for your second adjustment, which almost doubles from the first adjustment, it almost quadruples from the first -- from the original MSKCC agreement, the basis was a model that you used from Juno in 2016. I think you summarized that on an exhibit, correct? A. There is two parallel models from Juno. Q. Right. So you looked at an earlier one in 2016 and you compared it to a later one in 2017, right? A. It's, um, you know, early 2016 to just after the hypothetical negotiation, and it provides an apples-to-apples comparison, because they are the two similar financial models that were produced by Juno. Q. That spreadsheet is what you summarized on the left side of Exhibit 1247, correct? A. I don't have that committed to memory, but if we could pull it up, there is two general columns. Well, there is a first column that just lists the different labels, like what each row represents, but there is two primary columns of data, and those are reflects the early model versus the late

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model --1 2 Q. All right. 3 A. -- for Juno. And it reflects the differences in the marketplace at that point in time. 4 Q. Yeah. And you are relying on a note -- on a projection 5 in 2016 that indicated that Novartis had 10 percent of the 6 7 market, and you are comparing that to other projections a 8 year later, right? A. There are different market shares for different 9 10 indications. So if we look at the document, you will see it breaks it down by indication, and it's, you know, more of a 11 12 roughly two-year difference. Q. And you have -- you never talked to anyone at Juno about 13 the -- up until the time you were deposed in this case about 14 15 what the purpose of this model was, did you? A. I don't have that degree of specificity in terms of the 16 17 timeline. Q. You recall that at your deposition you were unable to 18 state what the purpose was of that model, right? 19 20 A. That doesn't immediately sound familiar. 21 Q. And you didn't speak to anyone at Juno about this model 2.2 before you relied upon it for your analysis, correct? 23 A. I do not recall speaking to folks at Juno about it. It was a subject of deposition testimony. So I could see what 24 they said about it under oath. But I do not recall offhand 25

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1	anyway discussions.
2	Q. Who testified about this model under oath, sir?
3	A. Um, I would have to take a look back at my report. I can
4	do that.
5	Q. Let's go back to the agreement you started with, which
6	was the MSKCC/Juno agreement. Under that agreement, am I
7	correct that Juno got not only patent rights but know-how?
8	A. Yes.
9	Q. And it got patent rights to more than the '190 patent,
10	correct?
11	A. That's right.
12	Q. And under the hypothetical negotiation, all Kite would
13	get is a bare license to the '190 patent, right?
14	A. That's right.
15	Q. And the know-how that Juno got but that Kite would not
16	get, you don't believe that was worthless, do you?
17	A. No, not worthless. It does not affect the royalty rate
18	under that agreement, which is 7.25 percent for just the
19	patent involved.
20	Q. Could you just answer my question, please, so we can move
21	on?
22	MR. WEINBERGER: I'm going to move to strike
23	everything after the first sentence, Your Honor.
24	THE COURT: So everything after "no, not worthless"
25	is struck. And the jury will be ordered to disregard it.

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Τ	BY MR. WEINBERGER:
2	Q. All right. So were you in court when Dr. Sadelain
3	testified the other day?
4	A. Yes, I was.
5	Q. Did you hear him say that this know-how was very valuable
6	to Juno?
7	A. Yes, to that effect, or at least that was part of the
8	takeaway.
9	Q. And you don't disagree with that, do you?
10	A. No, I don't.
11	Q. In fact, under the license agreement that you just
12	referred to, there is a provision that states that if in
13	substance, that if there is no valid patent on what was
14	transferred for MSK from MSKCC to Juno, a license fee of
15	3.625 percent would still be due, correct?
16	A. That's right.
17	Q. And Juno wouldn't have agreed to pay that for the
18	know-how if it thought it was not valuable, right?
19	A. I would think that know-how would be considered to be
20	valuable.
21	Q. But you didn't make any adjustment for the licensed
22	technology in the MSKCC Juno agreement compared to the
23	licensed technology at the hypothetical negotiation, right?
24	A. It would be inappropriate to make an adjustment for
25	know-how based upon the

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Q. Sir, did you make an adjustment or not? 1 2 A. No, I did not. 3 Q. Thank you. Now, let's talk a little about the first mover advantage. 4 Isn't it true that at the hypothetical negotiation 5 in October 2017, Juno expected to be third in the market 6 7 behind Novartis and behind Kite? 8 A. In effect, yes. Q. You say that in your report in paragraph 113, right? 9 10 A. I would imagine that I do. I don't recall the paragraph 11 number. 12 O. Okay. So at the hypothetical negotiation, Juno was not giving up the right to be first to market, right? 13 A. They were granting a first mover advantage to Kite and 14 15 that --Q. Sir, please, can you answer my question. Juno was not 16 17 giving up the right to be first to market; it had already 18 given that away, correct? 19 A. In effect. Q. And when I say given it away, I don't mean for free. 20 21 They gave it to Novartis in exchange for the license 2.2 agreement that we've talked about, right? 23 A. Yes, but KYMRIAH did not have --24 Q. Sir, please. Your counsel will have an opportunity. 25 Please just answer the question.

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And not only did Juno know that they would be third to market, they knew they would be third to market in the DLBCL indication that JCAR17 was supposed to target, right? A. There was an expectation that Novartis would get 3L DBCL (sic) approval shortly after the hypothetical negotiation in 2017. Q. Now, the second model that you used, the one you compared the 2016 model to to make this adjustment, that was a document that was referred to as Maple Company model. Do you remember that? A. I do. Q. And a particular part of the tab of the spreadsheet called JCAR17 assumptions was referenced in your attachment D4 to your report, correct? A. Possibly. Let me just take a quick look. Q. Take a look. A. Which tab? Q. This is a little difficult, because it's a -- it's a spreadsheet document, but you relied on, it's referenced in D4, row 359, and you got data from row 374 to 376. A. I'm sorry. I'm still not following your question. So I'm at D4, and I reference the Maple Company model, and I'm looking at a particular tab on that model relating to JCAR17. Q. Okay. Well, I'm going to put up a screenshot of the tab that you referenced, and I want to ask you a question about

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1	it. Can we do that?
2	MR. HEINRICH: This is not in evidence. We would
3	ask that it be taken down.
4	MR. WEINBERGER: All right.
5	BY MR. WEINBERGER:
6	Q. This is a spreadsheet that you relied on, which was a
7	Juno document, to make your adjustments, correct? This was a
8	spreadsheet you relied on?
9	MR. HEINRICH: Are you going to move it into
10	evidence or
11	MR. WEINBERGER: No, I'm trying to just lay a
12	foundation, and then I'll move it into evidence.
13	BY MR. WEINBERGER:
14	Q. You relied as we established in your report, D4, this
15	was the second set of projections that you relied on in
16	comparing when you compared the competitive impact from
17	2016 to 2017, right?
18	A. There is a particular financial model I rely upon, and
19	it's designated by what is called a Bates number on
20	attachment D4. I don't know if that is what you are putting
21	up on the screen or not.
22	Q. All right. So let me
23	THE COURT: I'm sorry. Can I see Mr. Chu and
24	Mr. Dane at sidebar very quickly, please.
25	(At the bench.)

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1	THE COURT: I'm just concerned about the time. I
2	know we have to get Mr. Schuetz on. And I don't want to
3	interrupt Mr. Weinberger. But maybe if we're going to
4	continue for a longer period of time, you may want to call
5	Mr. Schuetz out of order, if that's agreeable.
6	MR. CHU: That is agreeable to us.
7	MR. DANE: To interrupt the cross and resume it
8	later?
9	MR. CHU: Yes.
10	MR. DANE: When is Your Honor hoping to adjourn?
11	THE COURT: We have two jurors who are in one in
12	Lancaster, one juror in Lancaster, so that's a long drive to
13	go home. So I don't want to go beyond 4:15, 4:30.
14	MR. DANE: Okay. Let me can I talk to
15	Mr. Weinberger and Mr. Lawton?
16	THE COURT: Sure.
17	MR. CHU: Judge, while we are here, my apologies,
18	Your Honor, to you and everyone in the courtroom. I was
19	doing some work, and I didn't realize you anticipated
20	THE COURT: Don't worry about it.
21	MR. CHU: Thank you.
22	(In open court:)
23	MR. DANE: Your Honor, what Your Honor proposed we
24	are amenable to and to bring in Mr. Schuetz.
25	THE COURT: Is this a good time to

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1 MR. WEINBERGER: Yes. Absolutely, Your Honor. 2 THE COURT: There is a witness that needs to be 3 called out of order, so we are going to stop with the examination of Dr. Sullivan and move to the next witness. 4 Let's take a short recess before we do that, 5 6 10 minutes. During your absence do not discuss the case 7 amongst yourselves or any other person. I think we need some 8 air in here, I'm getting the impression. (Thereupon, the jury retired from the courtroom.) 9 10 THE COURT: Let's bring the jury in. 11 THE CLERK: All rise for the jury, please. 12 (Thereupon, the jury returned to the courtroom.) 13 THE COURT: Could we have all the jurors be 14 assembled with counsel. And the defendants are going to call 15 a witness. With the agreement of counsel for plaintiffs, a witness is being called out of order. 16 17 MR. LAWTON: Your Honor, Kite calls Dr. Thomas 18 Schuetz. THE CLERK: Please right here, doctor, come forward. 19 Would you please raise your right hand to be sworn. 20 21 THEREUPON: 2.2 THOMAS SCHUETZ, 23 called in these proceedings and being first duly sworn, 24 testifies as follows: 25