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                          UNITED STATES DISTRICT COURT
 2
                        FOR THE DISTRICT OF NEW JERSEY
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    HELSINN HEALTHCARE, S.A. and
    ROCHE PALO ALTO, LLC,
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                                        CIVIL ACTION NUMBER:
               Plaintiffs,
 6
                                               11-3962
                -vs-
 7
    DR. REDDY'S LABORATORIES, LTD.,
                                                TRIAL
    DR. REDDY'S LABORATORIES, INC.,
    TEVA PHARMACEUTICALS USA, INC.,
                                      WITH SEALED PORTIONS
    and TEVA PHARMACEUTICAL
    INDUSTRIES, LTD.
10
               Defendants.
11
         Clarkson S. Fisher United States Courthouse
12
         402 East State Street
         Trenton, New Jersey 08608
13
         June 15, 2015
14
    BEFORE:
                         THE HONORABLE MARY L. COOPER
                         UNITED STATES DISTRICT JUDGE
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    Certified as True and Correct as required by Title 28, U.S.C.,
    Section 753
24
    /S/ Regina A. Berenato-Tell, CCR, CRR, RMR, RPR
25
    /S/ Carol Farrell, CCR, CRR, RMR, CF
                                              Dr. Reddy's Laboratories, Ltd., et al.
                                                Helsinn Healthcare S.A., et al.
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11 S Patent No. 8 729 094

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25
    /S/ Carol Farrell, CCR, CRR, RMR, CCP, RPR, RSA
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1 THE COURT: -- item, which would be --2 THE WITNESS: Product specification, yeah. 3 THE COURT: And, so, although you use different 4 words, your slides, in terms of process definition, are not 5 that different, are they? How are they different? 6 THE WITNESS: No. I mean, I think the difference is 7 that the product profile in Dr. Kirsch's definition is 8 something that might -- like, you get a report and the 9 formulator is charged, okay, this is what we want to do; you 10 go make it. My position is the formulator is involved in the 11 product profile. 12 THE COURT: Okay. I thought I understood that to be 13 your point. 14 BY MR. DITTMANN: 15 And just if we can focus just on PDX 709 for one moment. 16 Just to make sure the record is clear, it's your experience 17 that the formulator team member of this development team would 18 be involved with all four steps seen on PDX 709, correct? 19 A . Yes. 20 Now, we have talked about the first two steps a bit 21 already in connection with your background. 22 With respect to this third step we see here, 23 considering the dose, volume, clinical parameters, can you 24 briefly describe how this work is typically accomplished in a



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drug development team?

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