

LSK Global PS <원격 모니터링 웨비나(Remote Monitoring Webinar)> 아젠다		
<b>2020. 12. 9 (수) 오후 3시 ~ 5시</b>		
<b>Session 1. Disruptive Innovation in Post-Covid19 Clinical Trials</b>	<b>LSK Global PS 이영작 대표이사</b>	<b>20 min</b>
<ul style="list-style-type: none"> <li>Why you need to remote monitoring?</li> </ul>		
<b>Session 2. Understanding E-Source in Clinical Trials : What you need to know</b>	<b>LSK PM Dept.</b>	<b>20 min</b>
<ul style="list-style-type: none"> <li>eSource, Why/ What?</li> <li>Common Misconceptions Surrounding E-Source Solutions</li> <li>The nature of e-source compliance</li> <li>Integration compatibility</li> <li>E-Source and CTMS do not meet the same needs</li> </ul>		
<b>Session 3. Remote Monitoring</b>	<b>LSK CTM HQs</b>	<b>20 min</b>
<ul style="list-style-type: none"> <li>Remote Monitoring of on-site activities</li> </ul>		
<b>Session 4. Target e*CTR</b>	<b>LSK CDM HQs</b>	<b>15 min</b>
<ul style="list-style-type: none"> <li>What is Target e*CTR?</li> <li>e*CTR workflow</li> <li>Registration</li> <li>Managing Accounts</li> <li>eSource</li> <li>View eSource in Target e*CRF</li> <li>Why should I use the e*CTR</li> </ul>		
<b>Session 5. Statistical monitoring in clinical trials</b>	<b>LSK STAT Dept.</b>	<b>15 min</b>
<ul style="list-style-type: none"> <li>Statistical monitoring in clinical trials</li> </ul>		
<b>Session 6. Risk-Based Monitoring vs. Remote Monitoring</b>	<b>LSK PM Dept.</b>	<b>15 min</b>
<ul style="list-style-type: none"> <li>Risk-Based Monitoring vs. Remote Monitoring</li> </ul>		
<b>Session 7. Veeva System</b>	<b>LSK PM Dept.</b>	<b>15 min</b>
<ul style="list-style-type: none"> <li>System instruction &amp; Demo</li> <li>CTMS : Study Dashboard / Report &amp; PD &amp; Issues</li> <li>eTMF : Essential Document viewer</li> <li>eISF : Source document review for remote SDV</li> </ul>		