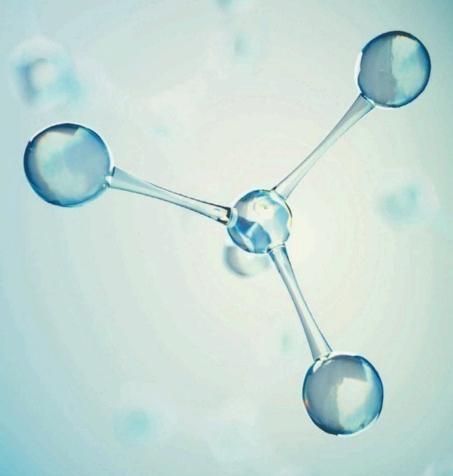
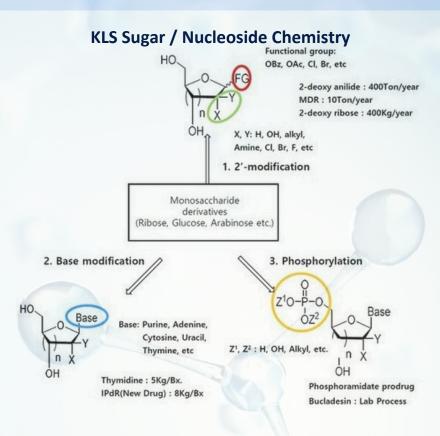
PHOSPHORAMIDITES



Summit Pharmaceuticals Europe

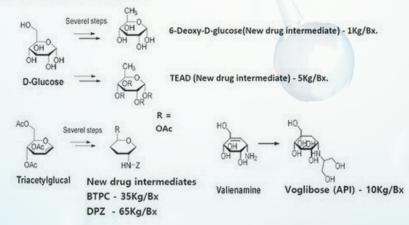
Overview

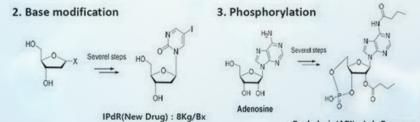
Kolon has been supplying **refined high-purity APIs and Intermediates** using our core technology, **Sugar Chemistry**. Our sugar chemistry consists of (1) control of functional groups (2) various refinement & crystallization methods (3) scale-up experience



New Drug Intermediate / API

1. 2'-Modification derivatives





Bucladesin(API): Lab Process

Product List

Kolon hascompleted the synthesis of **23 Phosphoramidites** through sugar / nucleoside chemistry. Each of them can incorporate flow chemistry, scale up, specific quality control and customization.

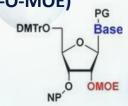
DNA Phosphoramidites

TBDMS-RNA Phosphoramidites

2'-Modified Phosphoramidites (2'-OMe)



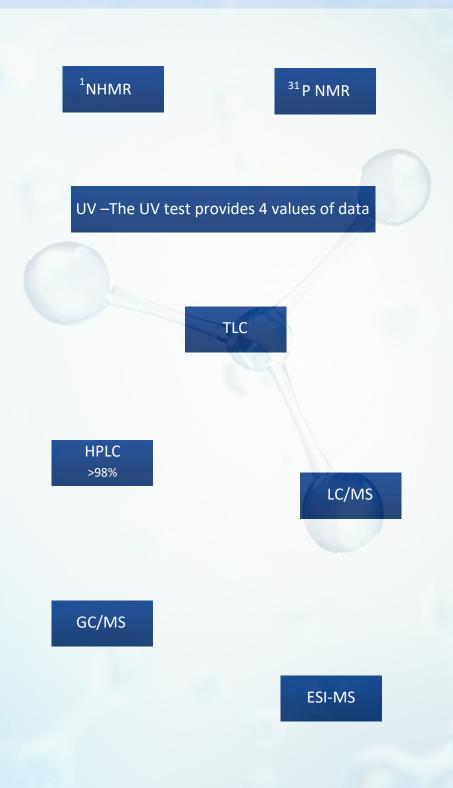
2'-Modified Phosphoramidites (2'-O-MOE)



2'-Modified Phosphoramidites (2'-F)

Quality Control

As the basic quality control methods for Phosphoramidites, Kolon conducts ¹H NMR, ³¹P NMR, UV, TLC,HPLC. LC/MS, GC/MS and MASS Spectrum are used for stricter management of impurities that has adverse effects on Oligo synthesis.



Differentiation

- Securing productcompetitiveness through differentiated process development
- Preoccupation of the market through early establishment of GMP process

Stage 1. Fast Follower

- ✓ Completion of general-purpose phosphoramidite development through competitor's product analysis
 - ·Continuously expanding the product portfolio with customer requests
- ✓ Promotion through collaboration with top 3 oligonucleotide makers around the world

Stage 2. Differentiation

- Securing cost competitiveness through differentiated process development
 - ·Reduce manufacturing costs by developing switchable continuous manufacturing process suitable for small-scale multi-variety production
 - ·Development of crystallization process to replace column process
- ✓ Diversification of Portfolio and development of customization
 - Development of a product line that can be used to manufacture oligonucleotides in the future (Diester, Triester, etc.)
 - •Development of various product lines to meet customer needs (Back-bone modification, 5'-capping, etc.)

Stage 3. First Mover

- ✓ It is scheduled to build a production facility of 300 kg per year in the Kilo-Lab Center in 2025
- ✓ A production base of 3 MT per year will be established in the GMP plant in 2027
 - •There is a high probability that regulation will be strengthened as the RNA market grows

