

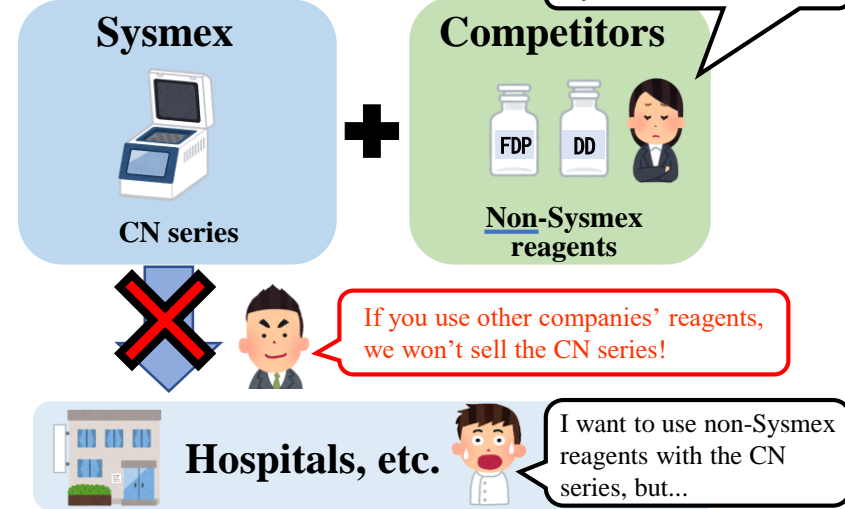
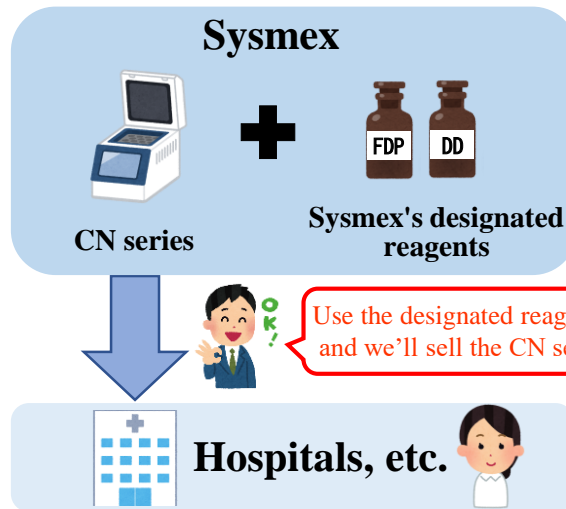
Overview of the Alleged Conduct

Tie-in Sales

Sysmex required hospitals and other medical institutions to use **only** its designated reagents when using specific blood coagulation analyzers (CN series) to measure “D-dimer” or “FDP,” even though reagents from other manufacturers could also be used.

Sysmex conditioned its supply of specific blood coagulation analyzers on the purchase of its designated reagents.

Sysmex is a leading company in the blood coagulation analyzer market.



There is a risk that suppliers selling non-Sysmex reagents may be excluded from transactions with hospitals or that their opportunities to trade with hospitals may decrease.

Overview of the Commitment Plan

1. Confirmation of the discontinuation of the alleged conduct, etc.

3. Refraining from engaging in similar conduct to the alleged conduct (five years) (This applies to all of Sysmex's blood coagulation analyzers and reagents as well.)

5. Establishing a contact point for handling inquiries from purchasing hospitals, etc., regarding conducts that potentially violate the Act.

2. Notifying hospitals, wholesalers, etc., and ensuring dissemination to employees, etc.

4. Creating codes of conduct, disseminating them to employees, etc., and conducting periodic training and audits.

6. Third party monitoring and reporting on each matter (five years)*

(Note) The item marked with an asterisk (*) is a measure based on the policy for more efficient and effective implementation of the commitment procedures, which was announced at the regular press conference of the Secretary General held on July 3, 2024.