## Study Request Form (SIAA Registration) (Example) Date:

		`	. Bute.					
Company			Study Monitor (name, position)					
Address		TEL	FAX					
Type of study								
	•		(Lot No.	)				
Control substance		(Lot No.						
Test substance	ce after study	Return / Storage						
completion								
Deadline of final report		Desired:						
	TEL FAX  The of study Acute oral toxicity/mutagenicity/mimary skin irritation/skin sensitization  st substance (Lot No. )  trol substance  stance after study ompletion (Lot No. )  of final report Desired:  of the chemical (by IUPAC nomenclature system, etc.):  Abbreviation:  ural, rational or molecular formula, molecular weight:  e enter detailed test substance information (composition, etc.)  ached sheet (if this column is insufficient)  or content (composition of active and other ingredients): ingredient xx wt%, other ingredients oo wt%  iption: viquid (powder) ility (Solubility/insolubility in water, organic solvents, etc.): soluble in XX, insoluble in OO (practically insoluble) ity (stability in solvents, to heat, to light, etc.; stability period): in the vehicle and to heat and light ge conditions (required if any): temperature (room temperature, under refrigeration, frozen)/light-resistant, hermetically closed							
<ol> <li>Structural, rational or molecular formula, molecular weight:         Please enter detailed test substance information (composition, etc.)     </li> <li>See attached sheet (if this column is insufficient)</li> </ol>								
Active ingredient xx wt%, other ingredients oo wt%  4. Description:								
	ubstance after study completion  "during the study materials storage period (fee-based); thereafter return or storage (fee-based)) ine of final report  Desired:  me of the chemical (by IUPAC nomenclature system, etc.):  Abbreviation: uctural, rational or molecular formula, molecular weight: ease enter detailed test substance information (composition, etc.)  attached sheet (if this column is insufficient)  ity or content (composition of active and other ingredients): tive ingredient xx wt%, other ingredients oo wt%  scription: low liquid (powder) ubility (Solubility/insolubility in water, organic solvents, etc.): ly soluble in XX, insoluble in OO (practically insoluble)  bility (stability in solvents, to heat, to light, etc.; stability period): le in the vehicle and to heat and light  rage conditions (required if any):							
_	•	•						
Freely solul	ble in XX, ins	soluble in OO (practically insolubl	e)					
•	•		ty period):					
7. Storage conditions (required if any): Storage temperature (room temperature, under refrigeration, frozen)/light-resistant, hermetically closed								
8. Precaution on handling (required if any):								
9. Other (designation of the test system, etc.):								

Notes

(1) Please let us know if you apply the test substance for approval.

(2) Regarding GLP studies, please let us know the results of pre- and post-study analyses and stability test of the test substance and stability in the solvents on procedural grounds.

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## Name and properties (purity, solubility, stability, etc.) of the test substance

Name of new chemical substance						
(by IUPAC nomenclature system)						
Alias name						
Structural or rational formula						
(if both are unclear, outline of						
manufacturing method)						
Purity of the new chemical substance	wt%		Lot No.			
Names of impurities and concentrations					wt	
CAS No.			Vapor pressure			
Molecular weight			Coefficient of partition			
Melting point	°C		Appearance at room			
Boiling point	°C		temperature			
Stability						
	Solvent	Solul	bility Stat		oility in the solvent	
	Water					
Solubility in solvents, etc.	DMSO					
	Ethanol					
	Other ( )					

[ Notes] Enter physicochemical properties to the extent possible as reference.

- 1. In the "Stability" column, enter stability to temperature, light, etc.
- 2. In the "Vapor pressure" column, enter vapor pressure of the test substance.
- 3. In the "coefficient of partition" column, enter coefficient of partition, temperature at measurement and the name of the solvent used for measurement.
- 4. In the "solubility in solvent, etc." column, enter solubility and stability of the test substance in the solvent.