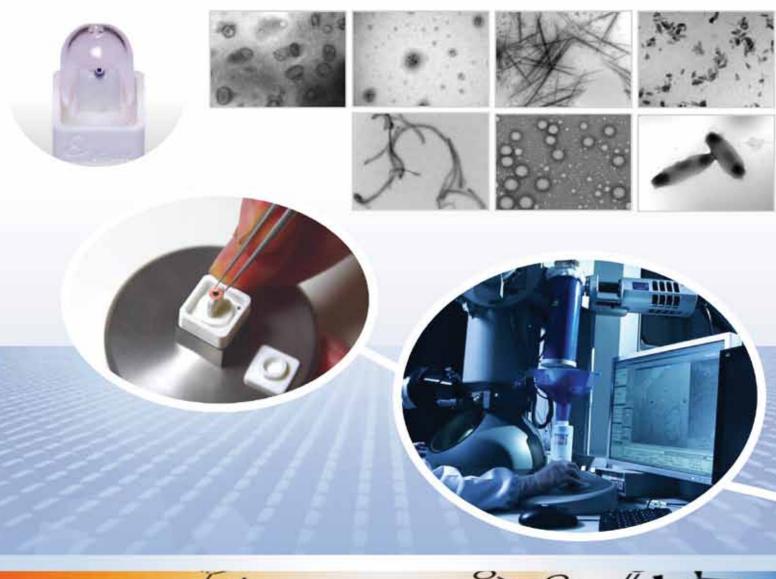
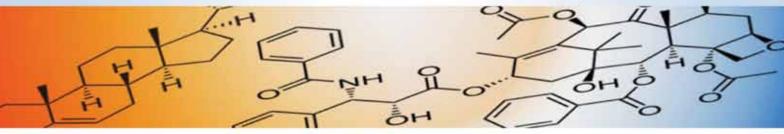
K-kit®





Innovative TEM specimen holder for liquid sample analysis







全国免费热线: 400 001 9621

商城:http://www.emcn.net.cn/

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MA-tek's enabling solution for liquid analysis by TEM



K-kit is a single-use sealable carrier with a microchannel inside. It's designed to facilitate convenient TEM and SEM observations of liquid samples, allowing nanoobjects, aggregates, and agglomerates (NOAAs) in liquid samples to be characterized.

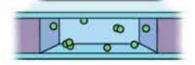






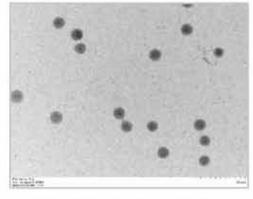
Transmission Electron Microscope (TEM)

Wet



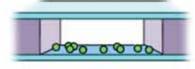
Filled with liquid

 The loaded liquid sample is sealed and imaged using TEM in the native liquid environment.



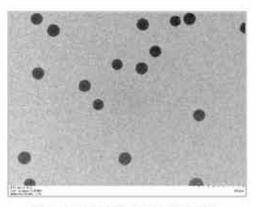
Acceptable image quality

Thin layer



Partially or fully dried

 A patented liquid drying protocol preserves the original morphology and physical state of nanomaterials with improved imaging resolution.

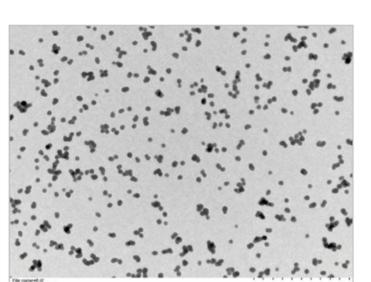


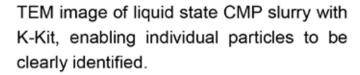
Very good image quality (NIST 100nm polystyrene spheres)



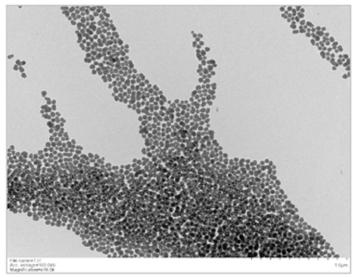
K-kit vs. TEM grid











TEM image of dried CMP slurry on Cu grid, unable to be analyzed individual particles due to agglomeration.

(\checkmark Good $\ \Delta$ Case dependent $\ X$ Not available)

Physicochemical Parameters	K-kit	Cu grid
1. Composition	✓	✓
2. Size	✓	✓
3. Shape	✓	✓
4. Size distribution	✓	Δ
5. Aggregation and agglomeration in liquid	✓	Х
6. Particle concentration	✓	Х
7. Liquid TEM observation	✓	Х

K-kit adaptability

Compatible with all kinds of TEM holders





If using some types of TEMs like JEOL 2100 which the z-axis focus depth less than ± 120 um, one will be possibly encountered the out-of-focus issue. In this situation, please refer to the possible solution as described on our website for it.

Good resistance to most solvents





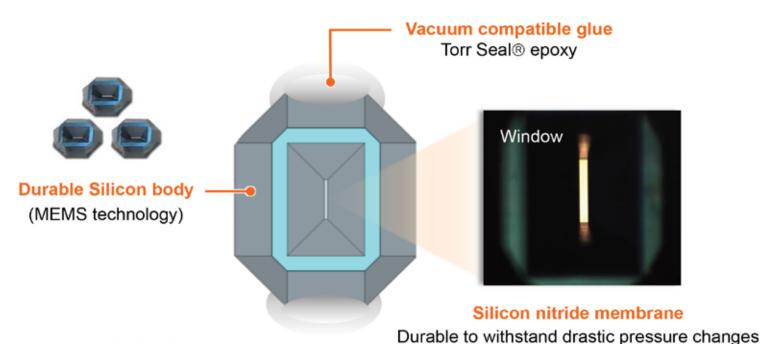
The following table shows the test results of Torr Seal Epoxy soaked in chemical solvents for 24 hours and then examined using FTIR (if dissolved) and visual observation (if dispersed).

	Water	PEG400	DMSO	Ethanol	0.1N HCI	0.1N KOH
	✓	✓	✓	✓	✓	✓
Compatibility	Toluene	NMP	ACN (CH ₃ CN)	Chloroform (CHCI ₃)	1% NH₄OH	0.1N HNO ₃
(FTIR)	✓	✓	✓	✓	✓	✓
	Hexane	IPA	Methanol	DCM	THF	Acetone
	√	✓	✓	A	A	•

Material and structural robustness

Broad temperature range for K-Kit -196°C to 120°C

Applicable with heating & cryo TEM holders





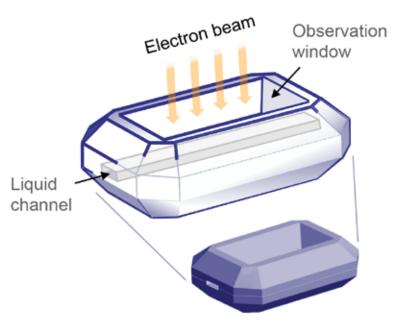
Channel tips

There are channel tips at both ends, to protect the surface condition and cleanness of the channel.



Reliable liquid loading

By capillary action, liquid can be loaded in a K-kit reliably, even the viscosity of it up to 3,000 mPa · s.



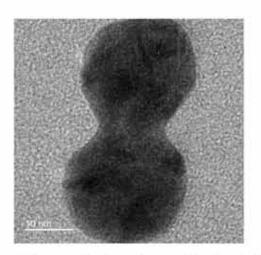
Unibody structure

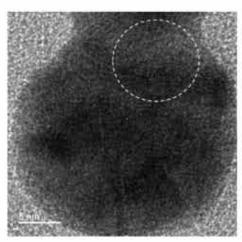
Cross-contamination free (Disposable), no need to do further assembly, surface treatment or pre-cleaning process before the use.

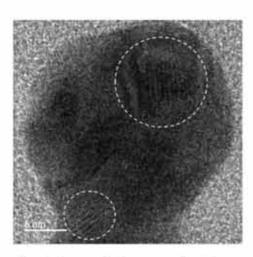
High image quality in TEM

Membrane	Sample Preparation			
Thickness of K-kit (Si ₃ N ₄)	Wet Mode	Dry Mode		
100nm	< 10nm	< 5nm		
30nm	< 5nm	< 2nm		

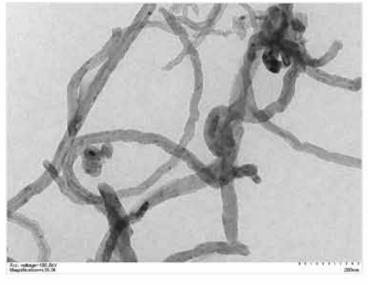


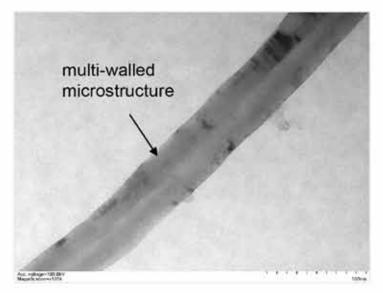






(Example) As shown in the TEM images of gold nanoparticles that formed from reduction process of AuCl₄ solution, the lattice lines of gold particles could be clearly observed by using Gap0.2um/ SiN30nm K-kit. (By FEI Talos TEM @ 200KV)



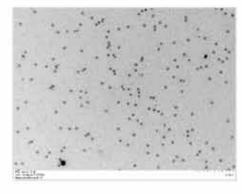


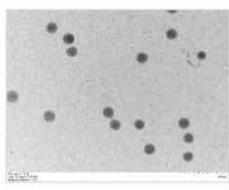
(Example) TEM images of multi-walled carbon nanotubes (WMCNT) that were fully dispersed in water. By using Gap2um/ SiN30nm K-kit, the structures of MWCNTs could be observed clearly. (By Hitachi HT7700 TEM @100KV; WMCNTs: OD 30-80 nm, Length <10 μm, 10wt%)

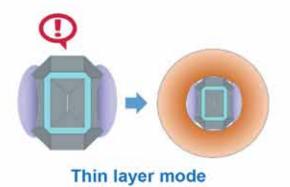
Wet and Thin Layer mode

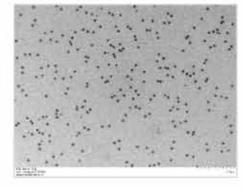
Sample preparation	Wet mode	Thin layer mode	
	With Liquid	Dry	
Imaging resolution	Good	Excellent	
Gap height suggested (μm)	0.1, 0.2, 0.5	0.5, 1, 2, 5	
Particle size (Loadable)	1nm~500nm	1nm~3000nm	
Particle shape	Keeping original	Potentially, could be deformed.	
Chemical reduction or potential damage by electron energy	High	Low	
Achievable	Fully filled	Thin liquid layer	
states of K-kit	Partially filled	Dry state	

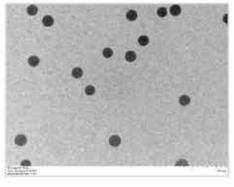






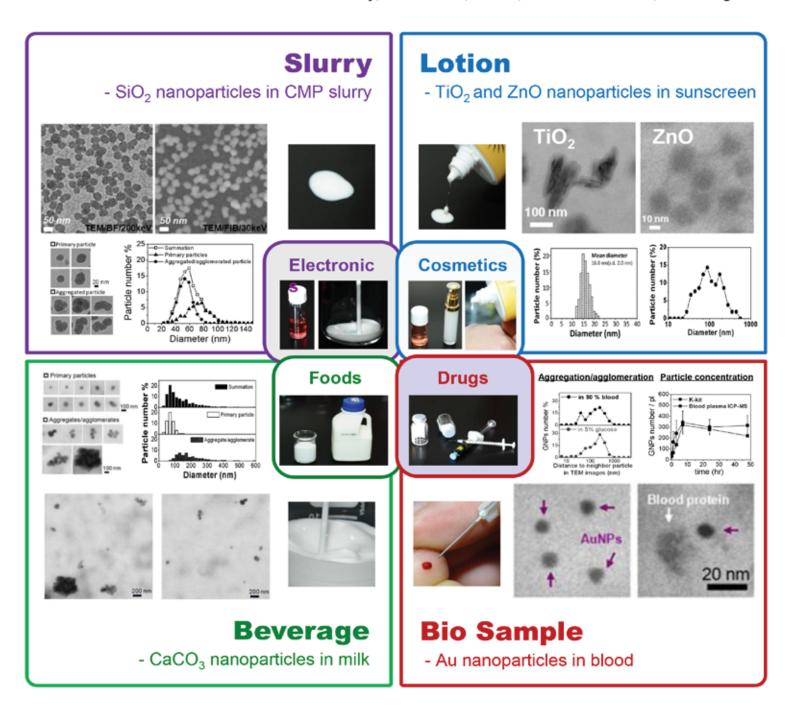






Application markets

Characterize NOAAs in electronics industry, cosmetics, foods, medical devices, and drugs.

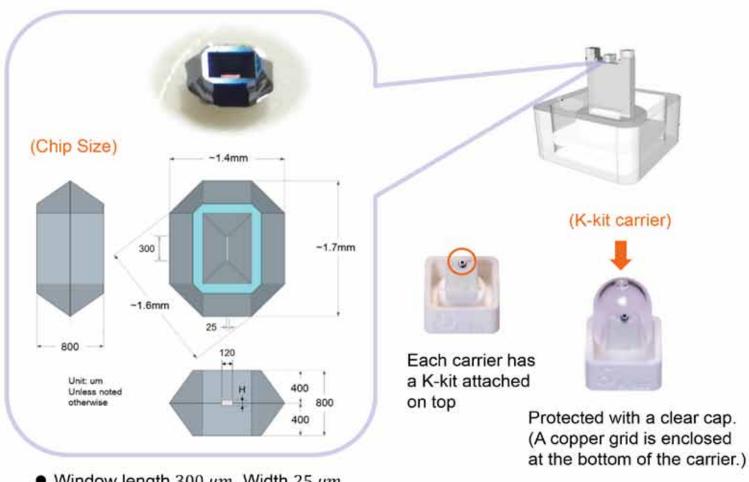


Reference:

- 1. US FDA 2012, Guidance for Industry Safety of Nanomaterials in Cosmetic Products.
- 2. EU/JRC July 2012, Requirements on Measurements for the Implementation of the European Commission Definition of the Term "Nanomaterials".
- 3. ISO/TR13014: 2012, Nanotechnologies -- Guidance on physico-chemical characterization of engineered nanoscale materials for toxicologic assessment.
- ICCR 2012, Characterization of Nanomaterials II Insolubility, Biopersistence and Size Measurement in Complex Media.



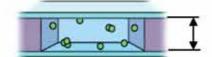
Shipping packages and tool sets



Window length 300 μm, Width 25 μm

K-kit

 Channel height (H): 0.2 and 2.0 standard 0.1, 0.5, 1.0 and 5.0 available



H = 0.1, 0.2, 0.5, 1,2, 5 (µm)



4 K-kits 6 K-kits

(Shipping packages)



Copper grid

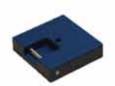


Channel opener



Starter box (glues, needles, channel opener, etc.)

(Tool sets and consumables)





Sample-loading K-kit gluing stand stage



Needle pen



K-kit tool box

Tool box, we offer a full tool set, including K-kit holder, sample-loading stage, needle pen, K-kit gluing stand, recommended glues, glass slides and some replacement parts.







Accessory Box



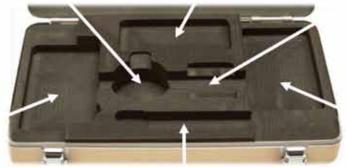
K-kit gluing stand



Glass-slide pack



Sample-loading stage





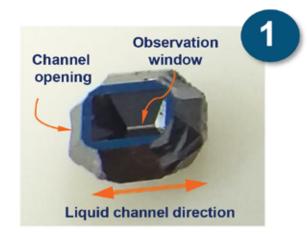
K-kit holder & needle pen



K-kit shipping package (without K-kits)



Sample preparation procedure



1.K-kit:

K-kits are Si-based microchannel devices with silicon nitride windows that allow SEM, FIB, STEM, and TEM observations. The shape is a result of anisotropic wet etching. The liquid channel is parallel to the window, with openings at both sides.



Liquid channel direction K-kit

2.Filling:

Liquid fills the channel through capillary force. The liquid surface is "pulled up" by the K-kit. Keep the K-kit steady for approximately 1 min to allow the filling to complete. The aqueous liquid sample should be placed on a glass slide. Both the K-kit and glass surface are hygroscopic. Do not immerse the K-kit in liquid to prevent from the window being contaminated.





3. Torr-seal:

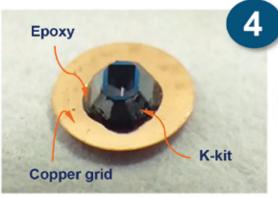
Cover the channel openings at both ends with Torr Seal epoxy after filling the device with liquid. (No need to do this gluing step, if one would like to dry out the liquid and leave the nanoparticles a Thin Layer mode in K-kit.)

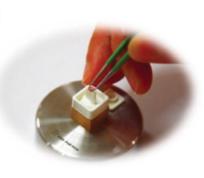


Use epoxy to mount the sealed K-kit to a copper grid by fitting it to the precut hole at the center of the grid.



QR code link to demo video





If on-line, please click the link to watch demo video: https://youtu.be/Hi9TyT4MwEg



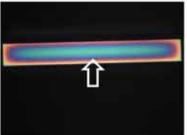


Matters needing attention when using K-kit

Inspection before use

- · With color rings on the membrane
- No any damage to the structure





With Newton's rings (Sealed by channel tips)

Channel tips removal

- · No color patterns on the film
- Liquid should be loaded in 30 minutes after removing the channel tips.

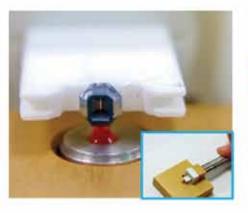


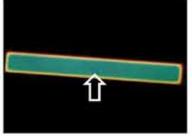


Flat membrane (Open to atmosphere)

Liquid loading

- Keep K-kit steadily touching on liquid surface at least for 30 seconds.
- Do not immerse K-kit in the liquid.

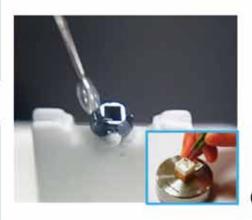


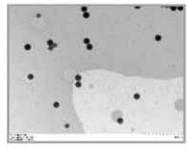


With color patterns (There's liquid filled)

Gluing process

- Glue the both ends of K-kit in 60 seconds after liquid loading.
- Do not glue off the K-kit, if making it with Dry mode.
- Glue with care, to avoid any adhesive flowing into the cavity.



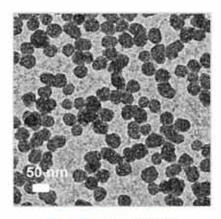


Liquid well reserved (Quickly to glue the K-kit)

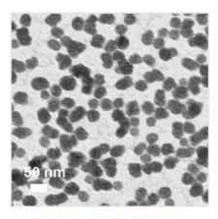


Available for TEM and SEM observations

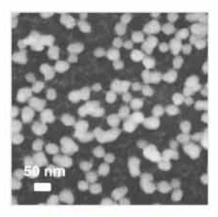
Compatible to versatile microscopy analyses







Hitachi-TEM @100Kev

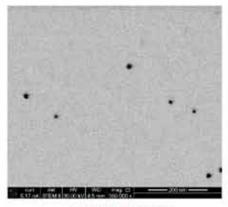


FEI-STEM @30Kev

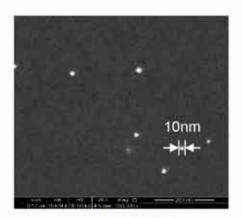
High imaging quality in SEM



Gold nanoparticles with sizes less than 10nm also could be clearly imaged with SEM.



Bright Field (BF)



Dark Field (DF)

☐ The comparison results of TEM and FIB-SEM images (Polystyrene beads in K-kit)

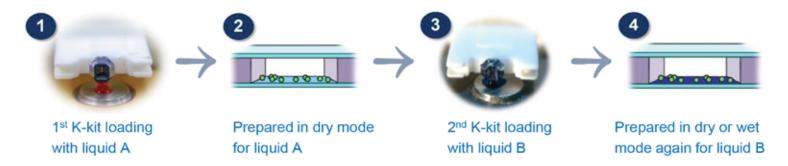
Hitachi HT7700 TEM FEI Helios 400 FIB-SEM DF

Multiple loadings and negative staining

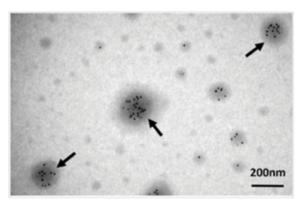
K-kit can provide researchers with faster and better choices when using a TEM to examine nanogranules of biological specimens in aqueous conditions.

Multiple loadings

With an unibody structure, K-kit can be used on multiple-loading applications, e.g. immunoelectron microscopy or catalyst chemistry studies etc.

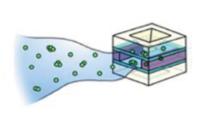


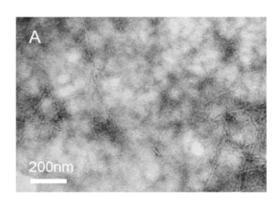
(Example) The presence of specific platelet granules could be labelled and observed by using a K-kit with multiple loadings. After some necessary pre-treatments and washings, the platelet sample in K-kit was incubated with the primary antibody (mouse monoclonal anti-P-selectin antibody) and next was reacted with a secondary antibody (6-nm gold-conjugated goat anti-mouse IgG antibody) for 2h at 37°C, and then the K-kit was sealed and examined in TEM. (Appl. Sci. 2020, 10, 4946)



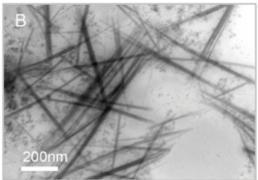
Immunoelectron micrographs of isolated platelet granules in a K-kit.

Negative staining





On Cu grid (In dry state)



In K-kit (With liquid).

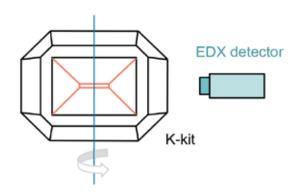
(Example) Negative staining TEM images of collagen on Cu grid and in K-kit. As shown in Fig. B, the collagen nanofibers could be clearly observed by using a wet-mode K-kit.

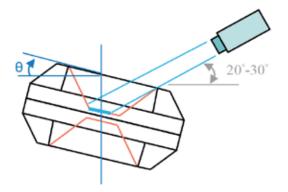


Available for EDX analysis

How to make EDX analysis achievable on a K-kit

In a TEM, an EDX detector is usually located at an angle of around 10 - 20° with regard
to the sample; X-ray signals excited from the observation window of a K-kit will be easily
blocked by that deep cavity. If so, by pointing the window long side of the K-kit toward the
EDX detector along with tilting the TEM holder at some angles over 15°, which can make
the EDX analysis achievable on it.



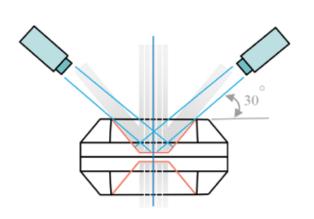


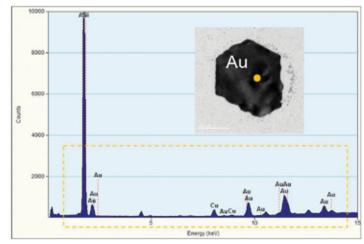
Align the window long side of K-kit to point to the EDX detector

Give a tilt at least of 15° toward the EDX detector

 For some modern TEMs that installed with multiple EDX detectors or only a detector but at a larger angle to the sample, one usually can get EDX signals from a K-kit directly without any tilting or rotation on the TEM holder.

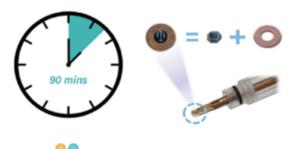






K-kit vs. in-situ TEM holder

K-kit can be the most convenient option in the market for liquid-TEM observation.



Around 90min required for 10 samples

Liquid loading and gluing for 10 K-kits (\sim 70min) + vacuum pumping (\sim 20min)





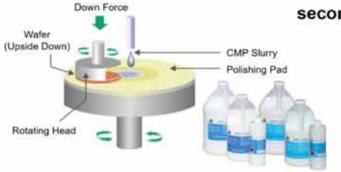
450min at least for 10 samples

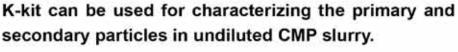
One by one; it needs the steps including surface treatment, assembly, leakage detection, and post-clean etc. for one sample. (> 45min for each)

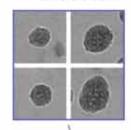
Product	K-kit	In-situ TEM holder
Cell size	1.7mm x 1.4mm (Fit in with Ø3 mm grids) > 2.4mm x 2.4mm	
Custom holder	No need	Required
Price	≤ US\$200	~ US\$100,000
Competitiveness	 Simple, quick and affordable Compatible to all TEM holders Available for SEM observation Good resistance to chemical solvents Cross-contamination free (Disposable) Achievable to quantitative analysis Reliable loading with viscous liquids Broad temperature range -196°C to 120°C 	Available for flowing and electrochemical studies
Weakness	 Only for static liquid analysis Electrodeless design 	 Sky-high prices Further pre-clean and assembly processes required With the risk of liquid leakage in TEM Dedicated for specific TEMs
User base	Industry and academia	Only for academia

NOAAs of abrasives in CMP slurry

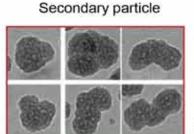
secondary particles in undiluted CMP slurry.





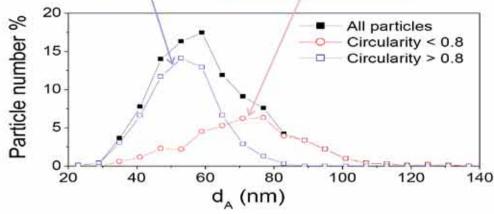


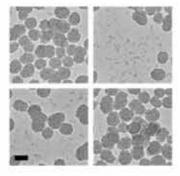
Primary particle



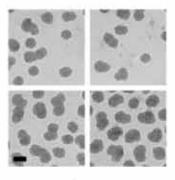
20 nm

- Composition
- Size/size distribution
- Shape
- Aggregation state
- Surface

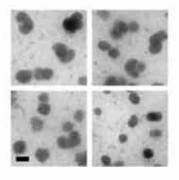




Dried on copper grid * Scale bar is 50 nm

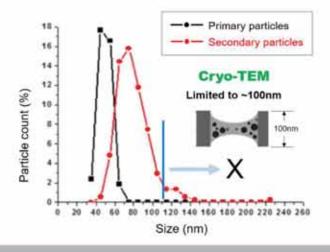


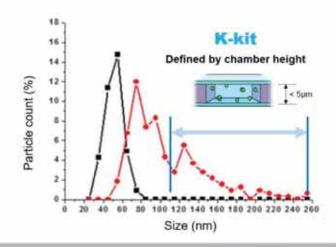
Frozen in Cryo-TEM grid



In liquid phase in K-kit

K-kit allows board size range of particles to be analyzed.

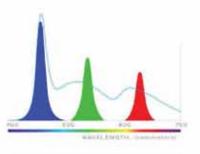


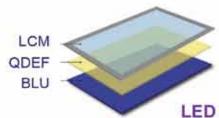




NOAAs of quantum dots in solution

Quantum dots will enable a market for devices and components over \$11bn by 2026.





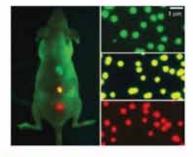
- QD: Quantum Dots
- · BLU: Backlight Unit
- LCM: Liquid Crystal Module
- QDEF: Quantum-dot Enhancement Film

LED display

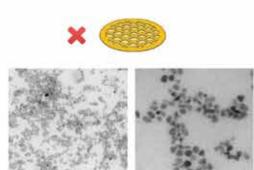




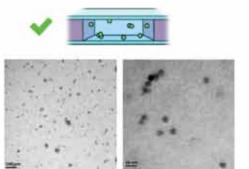




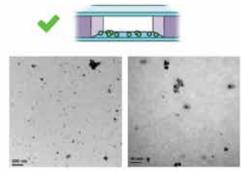
QD imaging diagnosis



QDs dried on copper grid

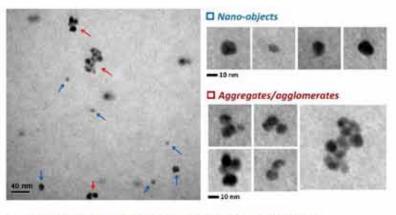


Wet mode of K-kit

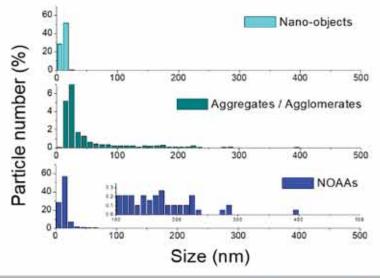


Thin Layer mode of K-kit

K-kit enabled, TEM images and size and size distribution of QDs in chloroform



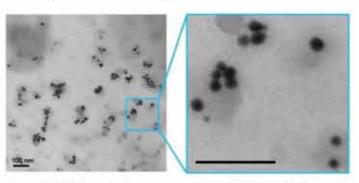
- · Sample solution was directly loaded into K-kit
- · Nano-objects = Primary particle
- Aggregates/agglomerates = Secondary particle



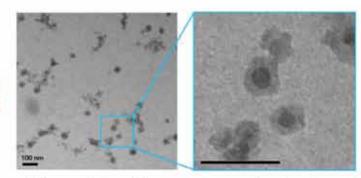


Drug particles in Nanopharmaceuticals

K-kit can be used for characterizing drug particles in Nanopharmaceuticals by imaging the particle morphology, size and size distribution, to evaluate drug formulation or conduct any bioequivalence study.



AuroVist® solution was directly loaded and sealed in a K-kit in liquid form.



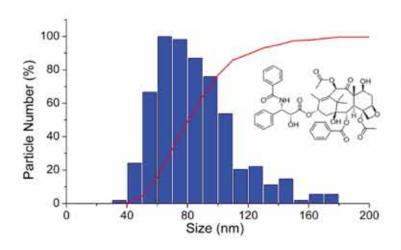
Oil emulsion in water was loaded and sealed in a K-kit in liquid form.

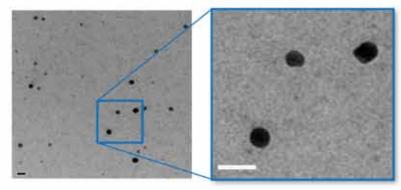
K-kit enabled, TEM images and size and size distribution of Abraxane in saline



Protein particles in Abraxane®

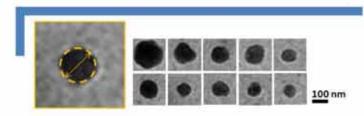
- Total calculated particle #: 319
- Average size: 85.1 nm
- Standard deviation: 27.0 nm





* Scale bar: 200 nm

Size/size distribution (D10, D50, D90)



Parameter	Size (nm)		
D 10	55.6		
D 50	80.1		
D 90	122.2		
Span: (D ₉₀ - D ₁₀) / D ₅₀	0.831		

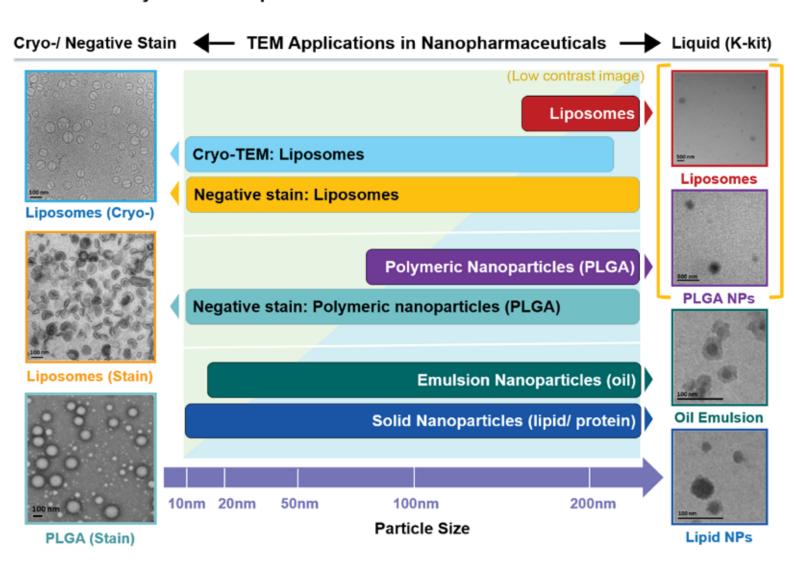
K-kit is the best option for Nanopharmaceuticals

◆ Applicable particle concentration for K-kit: 10¹¹~10¹⁴ particles/ml

The particle concentrations of most nano-drugs fall in the applicable range for K-kit. So, they can be directly observed and analyzed by using K-kit, without any dilution or condensation.

Brand Name of Pharmaceuticals	Doxil ® (1995 approved)	Abraxane ® (2005 approved)	Aurimune ® (Phase II)	Resovist ®	Rexin-G ® (Phase II)
Particle Size	80-100 nm	~ 130 nm	~ 27 nm (AuNPs core), ~ 30-40 nm as hydrated	~ 45-60 nm (Hydradynamic diameter)	~ 100 nm
Particle Concentrations	1.0 x 10 ¹⁴ liposome /ml	4.3 x 10 ¹³ albumin particles /ml	≤ 1.7 x 10 ¹² gold particles /ml	1 x 10 ¹⁴ particles /ml	1-4 x10 ¹¹ cfu

◆ The availability of K-kit compared with other solutions



Some bio-samples with very low contrast in TEM (like liposomes or PLGAs) that also can be clearly observed by using K-kits with negative staining.

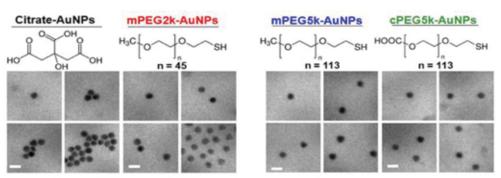
Quantitative characterization of nanoparticles in blood

Quantifying the concentration of nanoparticles in a biological matrix is important for in vivo analysis of their absorption, distribution, metabolism, and excretion, as well as for pharmacokinetic and toxicity studies. In this study, we demonstrated the possibility of using K-kit to obtain the aggregation and agglomeration states of nanomaterials in various native environments of interest.

(Tai et al. Anal. Chem. 2012, 84: 6312-6316)

Observation of different PEGylated gold nanoparticles in diluted blood with K-kits

	Par	Particle size		Surface properties		In 50 % Blood
	TEM ²	DL	S ^b	Zeta potential b	Surface ligand density c	Aggregation extents
Samples	d(nm)	d (nm)	PDI	$\xi(mV)$	PEG ($\#/nm^2$)	Aggregates (%)
Citrate-AuNPs	13.0±0.9	14.6	0.083	-28.3	non	87.1 ± 6.2
mPEG2k-AuNPs	27.5 ± 2.2	29.5	0.144	-23.0	2.27	28.4±9.2
mPEG5k-AuNPs	39.9 ± 2.8	39.6	0.071	-18.9	1.63	7.1 ± 3.9
cPEG5k-AuNPs	39.6±3.0	39.3	0.093	-35.5	0.82	17.3 ± 3.4



Citrate -GNPs

mPEG2k-GNPs

cPEG5k-GNPs

cPEG5k-GNPs

ion

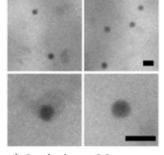
Distance to neighbor particle
in TEM images (nm)

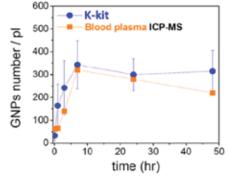
(TEM images of PEGylated gold nanoparticles in the 50% diluted blood)

The observation results demonstrated a high trend consistency on aggregation extent with the reference characterization information as listed in the table.

Quantification of the cPEG5k-GNP concentration in blood samples using K-kit and ICPMS analyses







* Scale bar: 20nm

This study showed the comparable results obtained for the number of cPEG5k-GNPs counted in the K-kit and measured by ICPMS. It confirmed that K-kit is a simple and convenient sampling device for evaluating the concentration of nanoparticles using TEM.

NOAAs of CaCO₃ NPs in milk

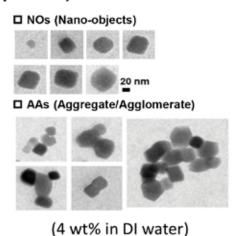


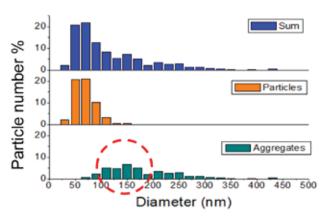
K-kit can be used for characterizing nanoobjects of foods in final product form, to evaluate the safety risks of nanomaterials in food additives and in substances in contact with foods.

Comprehensive physicochemical characterization								
P	arameter	Results	Methods					
1	Composition	Calcite CaCO ₃	TEM/EDX, XRD					
2	Size / size distribution	Average Diameter / Standard deviation						
	Crystal particle size	36 / 4 nm	XRD					
	Primary particle size	73 / 26 nm	TEM					
	Powder size	17 / 10 µm	SEM					
3	Shape	Cubic	TEM					
4	Aggregation/Agglomeration							
	in relevant media	Average diameter / Standard deviation	K-Kit / TEM					
	NOAAs	115 / 73 nm	(4wt% in DI water)					
	Nano-Objects	68 / 20 nm (number 58%)						
	Aggregations / Agglomerations	180 / 70 nm (number 42%)	42%)					
5	Solubility/Dispersibility	< 0.01% in Ca ²⁺ form	ICP/MS					
		Dispersed in DI water > 4 wt%	K-Kit/TEM					
		(20 ~ 450 nm)						
6	Surface charge	-23.4 ± 1.3 mV (in DI water)	Zeta potential					
7	Surface chemistry	Surface atom:	XPS					
	C (35%), O(48%), Ca(16%)							
8	Specific surface area	18.14 m ² /g	BET					

◆ Raw material (CaCO₃ powder)

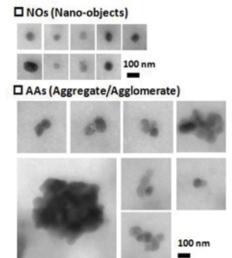


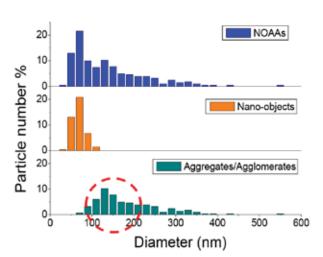




Final Product (Milk)







The aggregation extents of calcium granules might be slightly different between the raw additive and product form.

NOAAs of ZnO NPs in sunscreen

K-kit can be used for characterizing NOAAs of cosmetics in final product forms, including lotion, cream, and powder, to assess the safety risks of nanomaterials in cosmetic ingredients.

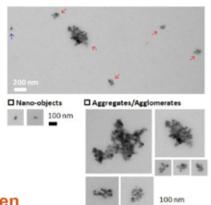


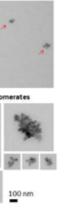
Regulations and regulatory trends for nanomaterials on cosmetics

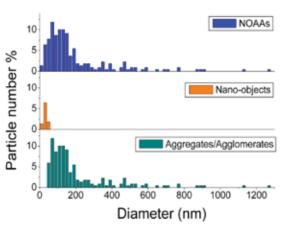
- International Cooperation on Cosmetic Regulation Report (ICCR) 2012 Characterization of Nanomaterials II – Insolubility, Biopersistence and Size Measurement in Complex Media.
- European Union (EU) Cosmetics Regulatory (EC) No. 1223/2009 Mandatory labeling of Nanomaterials as Ingredients in Cosmetics (Effective 2013/07/11)
- United States Food and Drug Administration Guidance (US FDA) 2012 Guidance of Industry - Safety of Nanomaterials in Cosmetic Products



ZnO NOAAs in sunscreen

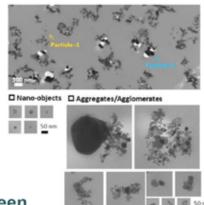


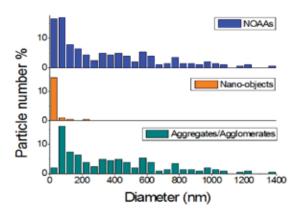




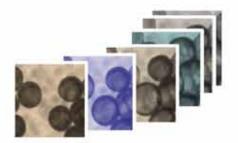


TiO₂ NOAAs in sunscreen



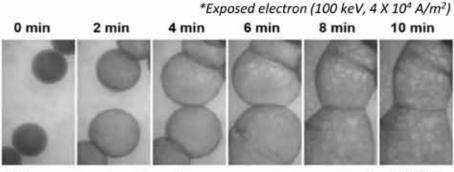


In-situ dynamic observations of NOAAs in liquid



The in-situ changes of nanomaterials can be observed and studied with K-kit dynamically, by a variation with time, area, temperature or surroundings.

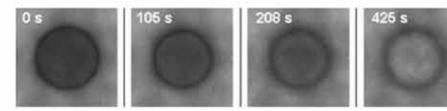




Silicate nanoparticles in water

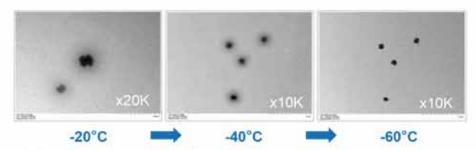






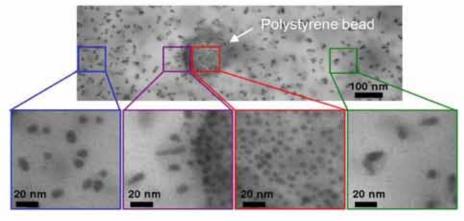
Polystyrene bead in PBS (Phosphate-buffered saline)





SiO₂ nanoparticles in Cryo-TEM





The growing Au nanoparticles nearby and far away from a polystyrene bead in AuCl₄ solution.



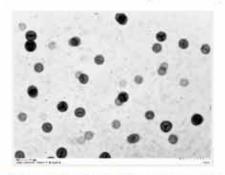
Enablement with K-Kit





Native state in liquid

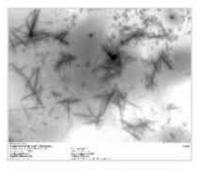
· Preserve the original morphology and physical state in liquid.



Extracellular vesicles of platelets



The nucleoid of E.coli



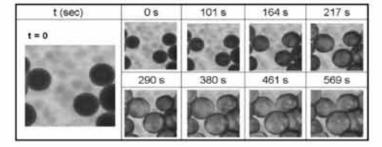
Collagen bundles in liquid



In-situ observation

 Kinetic mechanism of metal growth or physicochemical reaction process in liquid can be in-situ observed with increased reaction time.

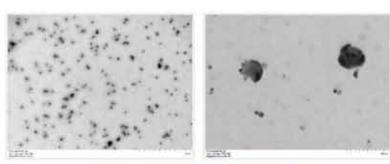
Dynamic observation of silicate nanoparticles



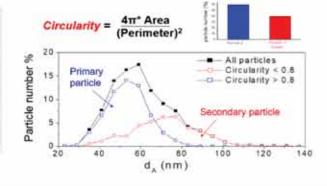


Quantitative analysis

Software of image recognition for nanoparticle size distribution analysis.



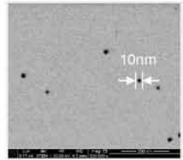
Nanoparticles of CMP slurry in K-kit

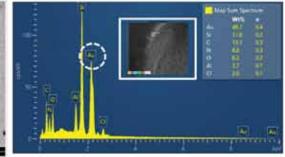




SEM & EDX analysis

 Nanoparticles in K-kit with sizes less than 10nm also could be nicely resolved in SEM.





Au nanoparticles were imaged and analyzed by FEI Helios 400 SEM

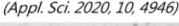


Enablement with K-Kit

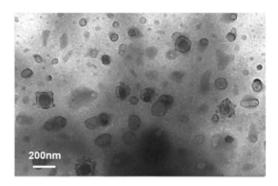




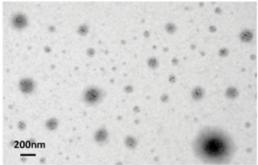
Negative staining and multiple loadings







Negative staining TEM image of isolated platelet granules in K-kit



Immunoelectron TEM image of platelet granules in K-kit (By multiple loadings)

- Some bio-samples which with very low image contrast in TEM can be also clearly observed by using K-kit with a negative staining treatment.
- With an unibody structure, it allows K-kit to be used on the applications with multiple loadings, e.g., immunoelectron microscopy studies.



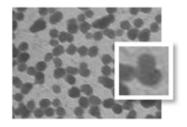
Wet and Dry modes

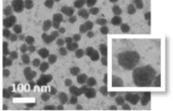
Wet mode:

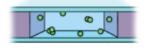
- With liquid fully or partially filled in K-kit.

Dry mode:

- A patented liquid drying protocol, with a thin liquid or fully dry state in K-kit. It can preserve the original morphology of nanomaterials along with the imaging results improved at the same time.







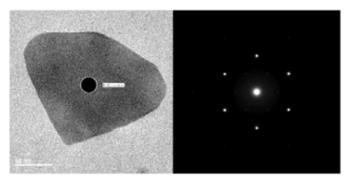


Undiluted Chemical-Mechanical Polishing (CMP) slurry directly loaded into K-kit.



TEM diffraction pattern

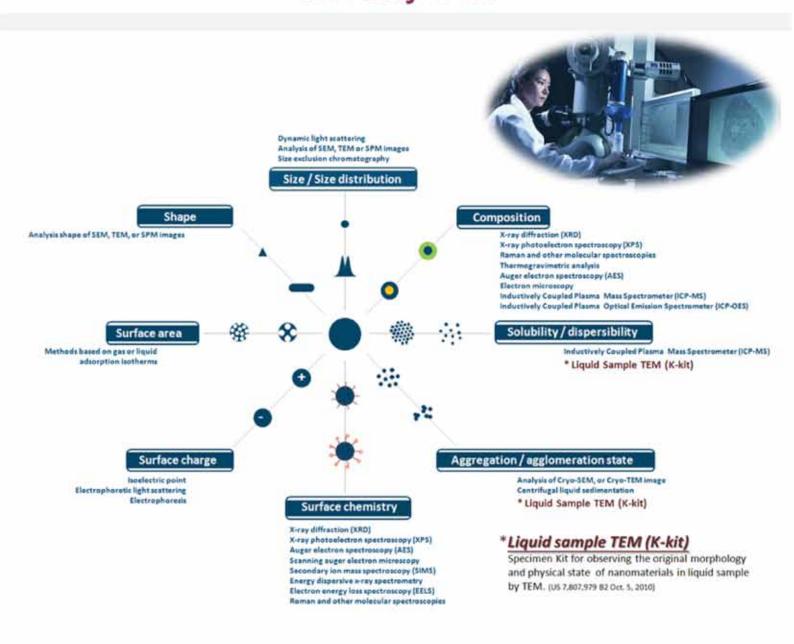
 TEM diffraction patterns of nanoparticles in liquid can be analyzed by using K-kit. In this example, Au nanoparticles were formed from reduction process with AuCl₄ solution and analyzed with FEI Tecnai TEM @200KV.

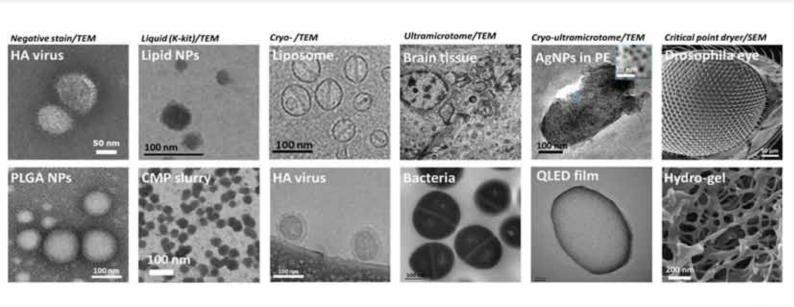


NBD (Nano-beam diffraction) result of a gold nanoparticle analyzed from a wet-mode K-kit.



Not only K-kit





Our services









Seven testing projects

- I. Analysis of the material's microscopic morphology
- II. Physical and chemical properties testing of materials
- III. Analysis of material composition
- IV. Elemental analysis
- V. Data Analysis
- VI. Sample Preparation (FIB. Ultrathin section, Ion thinning, etc.)
- VII. Environmental and Industrial Test

Service items

- SEM
- **XRD**
- **XPS**
- FT-IR
- Raman
- **ICP**
- MRI

- Organic Element Analysis
- Thermal Analysis
- **GPC**
- **AFM**
- **XRF**
- Particle Size and Zeta Potential
- MS
- 2D and 3D Analysis
- BET Hysteresis Loop

- Contact Angle
- **EBSD**
- Chemical Adsorption

About us

We are a professional testing institution in China. The test center aims to improve testing technology and take service research as its mission. Adhering to the concept of "integrity, seriousness, responsibility and efficiency". We provide scientific, accurate and reliable scientific research workers with professional perspective, advanced technology and excellent service. We have established long-term and stable cooperation with many well-known universities and research institutions and have been unanimously recognized.



Six advantages

Timeliness: Your samples will be immediately arranged after receiving, and the result of the regular samples will be delivered within 1-5 working days.

Professional: Our engineers are all researchers who have been engaged in many years of testing in the professional field and have rich experience.

Advantages of the instruments: Our center has complete facilities and has first-class testing instruments in the microscopic field.

Low Price: The charging standard is reasonable and fair, saving lot of funding for scientific researchers.

Guarantee: Perfect quality assurance system and after-sales service system to ensure the quality of test results.



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