

Supporting Information

Transfer and beyond: emerging strategies and trends in two-dimensional materials device fabrication

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Table S1. Scores of 6 evaluation criteria (ease of operation, cleanliness, transfer size, completeness, versatility, and multi-functionality) shown in the radar chart

Evaluation criterion		PMMA wet transfer method	Viscoelastic dry transfer method	Capillary force assisted hybrid method	Dielectric assisted method	SiN _x membranes pick up method	UV tape assisted hybrid method	Electrochemical stratification method	h-BN vdW pick up method	PC assisted transfer method
Ease of operation	Sacrificial layer	√	×	×	√	×	×	√	√	×
	tape	×	√	×	√	×	√	×	×	√
	Stamp	√	√	√	×	√	×	×	×	√
	Liquid solvent	√	×	×	√	√	√	√	√	×
	anneal	√	×	×	×	×	×	√	√	×
	Spending time	Hours	Minutes	Minutes	Minutes	Minutes	Hours	Minutes	1 Day	Half an hour
Cleanliness	Dielectric layer	PMMA	TRT/PDMS	PDMS	Sb ₂ O ₃ /PMMA/tape	SiN _x /Ta/Pt/Au	UV tape	PMMA	PDMS/PPC/h-BN	Glass/PDMS/PC/h-BN
	Evaluate	2	3	4	3	3	3	3	2	4
	Solvent residue	√	×	×	×	×	×	√	×	×
	Tape residue	×	√	×	×	×	√	×	×	×
	Scarified layer residue	√	√	×	√	×	×	√	×	×
	PDMS residue	×	√	×	×	×	×	×	×	×
Transfer size	Evaluate	2	3	5	4	5	4	3	3	4
	Scale	μm	μm		Wafer-scale	μm	Wafer-scale	Wafer-scale	μm	μm

Completeness	Evaluate	4	3	4	5	3	5	5	2	3
	wrinkle	√	×	×	×	×	×	√	×	×
	bubble	√	×	×	×	×	×	√	√	√
	breakage	√	√	×	√	×	×	√	×	×
Versatility	Evaluate	2	3	4	3	4	5	2	3	3
	CVD 2D materials	√	×	√	√	√	√	√	√	√
	Exfoliated 2D materials	√	√	√	√	√	√	√	√	√
	Evaporated/deposited thin film	√	×	√	×	√	√	√	√	√
Multi-functionality	Transfer of Electrodes	√	√	√	×	×	√	√	√	√
	Evaluate	4	2	4	2	3	4	4	4	4
	The operation accuracy is not high, which makes it difficult to apply techniques such as twist.	Some additional materials are needed to enhance adhesion and expand the functionality.	It can provide a high-quality interface and has excellent scalability.	It has a high-quality interface and can be patterned on dielectric films to achieve pattern transfer.	It has a high-quality interface, but there are certain requirements for the roughness of the substrate.	While providing a high-quality interface, it also has excellent scalability.	Can be combined with subsequent large-scale applications.	It can achieve high-quality interfaces, and h-BN can be used as the encapsulation layer.	Cannot completely control the existence of bubbles.	
	Controlled Strain, controlled wrinkle, ease for twist easy integration with other structures, etc.									
	Evaluate	2	2	4	3	3	4	3	4	3

Ref.			Ref ¹	Ref ²	Ref ³	Ref ⁴	Ref ⁵	Ref ⁶	Ref ⁷	Ref ⁸	Ref ⁹
Table S2.											
Year	Transfer method	Device structure	Contaminants	Carrier mobility		Contact resistance (Ω) and Threshold voltage (V)	Subthreshold swing (SS)	Long-term cycling stability	Residue removing strategies	Ref	
				Before treatment	After treatment						
2011	PMMA wet transfer (Wet) ¹⁰	graphene FET	PMMA	1400 cm ² ·V ⁻¹ ·s ⁻¹ ,	2700 cm ² ·V ⁻¹ ·s ⁻¹ ,	NA	NA	NA	Vacuum annealing	¹⁰	
2012	PMMA wet transfer (Wet) ¹¹	graphene FET	PMMA	1810 ± 710 cm ² ·V ⁻¹ ·s ⁻¹ ,	3200 ± 670 cm ² ·V ⁻¹ ·s ⁻¹ ,	500-750 Ω	NA	NA	300°C UHV anneal, and the in situ 80°C vacuum anneal	¹¹	
2016	Tape exfoliation (Dry) ¹²	MoS ₂ FET	Tape residue	NA	35 cm ² ·V ⁻¹ ·s ⁻¹ ,	740 Ω · μ m and -16V	NA	4-month air-stable	Vacuum annealing	¹²	
2016	PMMA wet transfer CVD grown graphene (Wet) ¹³	graphene FET	PMMA	Electron and hole mobility were 2141 and 2230 cm ² ·V ⁻¹ ·s ⁻¹ ,	electron and hole mobility were 3770 and 4232 cm ² ·V ⁻¹ ·s ⁻¹	557.3 Ω to 125.4 Ω	NA	NA	laser exposure	¹³	
2016	PVA-assisted transfer vs PMMA-assisted transfer (Wet) ¹⁴	graphene FET	NA	PMMA assisted: electron and hole mobility were 1607 and	PVA assisted: electron and hole mobility were 3587 and 3800 cm ² ·V ⁻¹ ·s ⁻¹	NA	NA	NA	DI water	¹⁴	

				1687 cm ² ·V ⁻¹ ·s ⁻¹ ,							
				PMMA assisted: electron were 1.1 cm ² ·V ⁻¹ ·s ⁻¹ ,	PVA assisted: electron were 12 cm ² ·V ⁻¹ ·s ⁻¹	-6.5v	30mV/dec to 22mV/dec	NA			
2016	PVP+PVA-assisted transfer (Dry) ¹⁵	MoS ₂ FET	N/A	PMMA assisted: 3.0 cm ² ·V ⁻¹ ·s ⁻¹ ,	PVA assisted: 9.0 cm ² ·V ⁻¹ ·s ⁻¹	NA	NA	NA	Vacuum annealing	¹⁵	
2016	PPC/PDMS stamp assisted transfer (Dry) ¹⁶	MoS ₂ FET	Polymer residue	NA	25 cm ² ·V ⁻¹ ·s ⁻¹	75 Ω μm	120 mV/dec	NA	400°C annealing	¹⁶	
2017	PMMA wet transfer CVD grown graphene (Wet) ¹⁷	graphene FET	PMMA	electron and hole mobility were 2047.1 and 1440.9 cm ² ·V ⁻¹ ·s ⁻¹ ,	electron and hole mobility were 2257.5 and 1500.3 cm ² ·V ⁻¹ ·s ⁻¹	NA	NA	NA	AFM cleaning	¹⁷	
2019	gel tape exfoliation (Dry) ¹⁸	MoS ₂ FET	photoresist	~6.5 cm ² ·V ⁻¹ ·s ⁻¹ ,	~100 cm ² ·V ⁻¹ ·s ⁻¹ ,	~20.2 kΩ·μm to ~1 kΩ·μm	60mV/dec	NA	O ₂ plasma	¹⁸	
2019	Scotch tape method (CVD-grown WSe ₂ , and	WSe ₂ FET	PMMA	32.5 ± 14.8 cm ² ·V ⁻¹ ·s ⁻¹ ,	36.7 ± 16.5 cm ² ·V ⁻¹ ·s ⁻¹ ,	+23.2V ± 1.3 to +20.2V ± 1.2	NA	NA	contact mode atomic force microscopy	¹⁹	

	exfoliated MoS ₂ and WSe ₂) (Wet) ¹⁹	MoS ₂ FET		98.5 ± 17.9 cm ² ·V ⁻¹ ·s ⁻¹ ,	100.6 ± 22.5 cm ² ·V ⁻¹ ·s ⁻¹ ,	-3.4V ± 5.5 to -6.1V ± 4.8	NA	NA	(AFM)	
2023	Ice-aided transfer (IAT) and ice stamp transfer (IST) (Hybrid) ²⁰	monolayer MoS ₂ FET	NA	NA	24.3 cm ² ·V ⁻¹ ·s ⁻¹	NA	NA	6-month air stable	Ice-assisted transfer	²⁰
2024	PPC assisted transfer (Wet) ²¹	monolayer MoS ₂ FET	0.08% PPC residue coverage	NA	19.2 cm ² ·V ⁻¹ ·s ⁻¹	78 Ω·μm and 5V	NA	NA	NA	²¹
2025	Capillary force-assisted PDMS stamp transfer (Hybrid) ²²	MoS ₂ FET	NA	NA	15.62 cm ² ·V ⁻¹ ·s ⁻¹	-0.52V	150 mV/dec	NA	NA	²²
2025	Hydrogenated diamond assisted transfer (Dry) ²³	monolayer MoS ₂ FET	PDMS	NA	45 cm ² ·V ⁻¹ ·s ⁻¹	3V	NA	Minimal degradation (less than 10%) over a 3-day period of measurements	300°C annealing for 1h to remove PDMS residue	²³

Reference:

1. A. Reina, H. Son, L. Jiao, B. Fan, M. S. Dresselhaus, Z. Liu and J. Kong, *J. Phys. Chem. C.*, 2008, **112**, 17741-17744.
2. A. Castellanos-Gomez, M. Buscema, R. Molenaar, V. Singh, L. Janssen, H. S. J. van der Zant and G. A. Steele, *2D Mater.*, 2014, **1**, 011002.
3. X. Ma, Q. Liu, D. Xu, Y. Zhu, S. Kim, Y. Cui, L. Zhong and M. Liu, *Nano Lett.*, 2017, **17**, 6961-6967.
4. J. Liao, Y. Zhao, X. Chen, Z. Hu, S. Bu, Y. Zhu, Q. Lu, M. Shang, H. Wu, F. Li, Z. Shi, Q. Zhao, K. Jia, J. Hu, Z. Han, Q. Xie, X. Zhao, J. Yin, W. Wang, H. Peng, X. Qiu, Y. Zhang, L. Lin and Z. Liu, *Nat. Electron.*, 2025, DOI: 10.1038/s41928-025-01353-x.

5. W. Wang, N. Clark, M. Hamer, A. Carl, E. Tovari, S. Sullivan-Allsop, E. Tillotson, Y. Gao, H. de Latour, F. Selles, J. Howarth, E. G. Castanon, M. Zhou, H. Bai, X. Li, A. Weston, K. Watanabe, T. Taniguchi, C. Mattevi, T. H. Bointon, P. V. Wiper, A. J. Strudwick, L. A. Ponomarenko, A. V. Kretinin, S. J. Haigh, A. Summerfield and R. Gorbachev, *Nat. Electron.*, 2023, **6**, 981-990.
6. M. Nakatani, S. Fukamachi, P. Solís-Fernández, S. Honda, K. Kawahara, Y. Tsuji, Y. Sumiya, M. Kuroki, K. Li, Q. Liu, Y.-C. Lin, A. Uchida, S. Oyama, H. G. Ji, K. Okada, K. Suenaga, Y. Kawano, K. Yoshizawa, A. Yasui and H. Ago, *Nat. Electron.*, 2024, **7**, 119-130.
7. Y. Wang, Y. Zheng, X. Xu, E. Dubuisson, Q. Bao, J. Lu and K. P. Loh, *ACS Nano*, 2011, **5**, 9927-9933.
8. L. Wang, I. Meric, P. Y. Huang, Q. Gao, Y. Gao, H. Tran, T. Taniguchi, K. Watanabe, L. M. Campos, D. A. Muller, J. Guo, P. Kim, J. Hone, K. L. Shepard and C. R. Dean, *Science*, 2013, **342**, 614-617.
9. P. J. Zomer, M. H. D. Guimarães, J. C. Brant, N. Tombros and B. J. van Wees, *Appl. Phys. Lett.*, 2014, **105**, 013101.
10. A. Pirkle, J. Chan, A. Venugopal, D. Hinojos, C. W. Magnuson, S. McDonnell, L. Colombo, E. M. Vogel, R. S. Ruoff and R. M. Wallace, *Appl. Phys. Lett.*, 2011, **99**, 122108.
11. J. Chan, A. Venugopal, A. Pirkle, S. McDonnell, D. Hinojos, C. W. Magnuson, R. S. Ruoff, L. Colombo, R. M. Wallace and E. M. Vogel, *ACS Nano*, 2012, **6**, 3224-3229.
12. C. D. English, G. Shine, V. E. Dorgan, K. C. Saraswat and E. Pop, *Nano Lett.*, 2016, **16**, 3824-3830.
13. Y. Jia, X. Gong, P. Peng, Z. Wang, Z. Tian, L. Ren, Y. Fu and H. Zhang, *Nano-Micro Lett.*, 2016, **8**, 336-346.
14. H. Van Ngoc, Y. Qian, S. K. Han and D. J. Kang, *Sci. Rep.*, 2016, **6**, 33096.
15. Z. Lu, L. Sun, G. Xu, J. Zheng, Q. Zhang, J. Wang and L. Jiao, *ACS Nano*, 2016, **10**, 5237-5242.
16. A. Nourbakhsh, A. Zubair, R. N. Sajjad, A. Tavakkoli K. G, W. Chen, S. Fang, X. Ling, J. Kong, M. S. Dresselhaus, E. Kaxiras, K. K. Berggren, D. Antoniadis and T. Palacios, *Nano Lett.*, 2016, **16**, 7798-7806.
17. W. Choi, M. A. Shehzad, S. Park and Y. Seo, *RSC Adv.*, 2017, **7**, 6943-6949.
18. P. Bolshakov, C. M. Smyth, A. Khosravi, P. Zhao, P. K. Hurley, C. L. Hinkle, R. M. Wallace and C. D. Young, *ACS Appl. Electron. Mater.*, 2019, **1**, 210-219.
19. J. Liang, K. Xu, B. Toncini, B. Bersch, B. Jariwala, Y.-C. Lin, J. Robinson and S. K. Fullerton-Shirey, *Adv. Mater. Interfaces*, 2019, **6**, 1801321.
20. H. Liu, Q. H. Thi, P. Man, X. Chen, T. Chen, L. W. Wong, S. Jiang, L. Huang, T. Yang, K. H. Leung, T. T. Leung, S. Gao, H. Chen, C.-S. Lee, M. Kan, J. Zhao, Q. Deng and T. H. Ly, *Adv. Mater.* 2023, **35**, 2210503.
21. A. Mondal, C. Biswas, S. Park, W. Cha, S.-H. Kang, M. Yoon, S. H. Choi, K. K. Kim and Y. H. Lee, *Nat. Nanotechnol.*, 2024, **19**, 34-43.
22. L. Liu, Z. Cai, S. Xue, H. Huang, S. Chen, S. Gou, Z. Zhang, Y. Guo, Y. Yao, W. Bao and P. Zhou, *Nat. Electron.*, 2025, **8**, 135-146.

23. K. Xing, D. McEwen, Y. Yin, W. Zhao, A. Bake, D. Cortie, J. Liu, T.-H.-Y. Vu, Y.-H. Chen, J. Hone, A. Stacey, M. T. Edmonds, N. V. Medhekar, K. Watanabe, T. Taniguchi, Q. Ou, D.-C. Qi and M. S. Fuhrer, *ACS Nano*, 2025, **19**, 3579-3588.