

IPC-MS-810

Guidelines for High Volume Microsection

Developed by the Automatic Microsectioning Task Group (7-10b) of the Testing Committee (7-10) of IPC

Users of this standard are encouraged to participate in the development of future revisions.

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Table of Contents

1	SCOPE
2 2.1	APPLICABLE DOCUMENTS
3 3.1 3.2	SAMPLE REMOVAL PROCESS 1 Sample Location 1 Removal Method 1
4 4.1	MOUNT PROCESS
4.2	Tooling System
4.3	Mounting 4
4.4	Mount Process Quality 5
5	GRIND PROCESS
5.1	Equipment 5
5.2	Tooling
5.3 5.4	Tool Stops
5.4 5.5	Consumables
	- ·
6	POLISH PROCESS
6.1	Equipment
6.2 6.3	Tooling
0.3 6.4	Consumables
6.5	Polish Process Quality
0.5	Tonsh Trocess Quanty
_	
7	MICRO-ETCHING
7.1	Application Methods 13
7.1 7.2	Application Methods13Types13
7.1	Application Methods13Types13TROUBLESHOOT GUIDE13
7.1 7.2 8 9	Application Methods 13 Types 13 TROUBLESHOOT GUIDE 13 GLOSSARY 14
7.1 7.2 8 9 9.1	Application Methods13Types13TROUBLESHOOT GUIDE13GLOSSARY14Abrasion14
7.1 7.2 8 9 9.1 9.2	Application Methods13Types13TROUBLESHOOT GUIDE13GLOSSARY14Abrasion14Charging14
7.1 7.2 8 9 9.1 9.2 9.3	Application Methods13Types.13TROUBLESHOOT GUIDE13GLOSSARY14Abrasion14Charging14Coupon Test Strip.14
7.1 7.2 8 9 9.1 9.2 9.3 9.4	Application Methods13Types13TROUBLESHOOT GUIDE13GLOSSARY14Abrasion14Charging14Coupon Test Strip14Crescent Moon Scratch14
7.1 7.2 8 9 9.1 9.2 9.3 9.4 9.5	Application Methods13Types13TROUBLESHOOT GUIDE13GLOSSARY14Abrasion14Charging14Coupon Test Strip14Crescent Moon Scratch14Grind14
7.1 7.2 8 9 9.1 9.2 9.3 9.3 9.4 9.5 9.6	Application Methods13Types.13TROUBLESHOOT GUIDE13GLOSSARY14Abrasion14Charging14Coupon Test Strip.14Crescent Moon Scratch14Grind.14Grinding Wheel14
7.1 7.2 8 9.1 9.2 9.3 9.3 9.4 9.5 9.6 9.7	Application Methods13Types13TROUBLESHOOT GUIDE13GLOSSARY14Abrasion14Charging14Coupon Test Strip14Crescent Moon Scratch14Grind14Grind Wheel14Grit Size14
7.1 7.2 8 9.1 9.2 9.3 9.4 9.5 9.6 9.7 9.8	Application Methods13Types13TROUBLESHOOT GUIDE13GLOSSARY14Abrasion14Charging14Coupon Test Strip14Crescent Moon Scratch14Grind14Grind Wheel14Grit Size14Grind Mount Holder14
7.1 7.2 8 9 9.1 9.2 9.3 9.4 9.5 9.6 9.7 9.8 9.9	Application Methods13Types.13TROUBLESHOOT GUIDE13GLOSSARY14Abrasion14Charging14Coupon Test Strip.14Crescent Moon Scratch14Grind.14Grinding Wheel14Grind Mount Holder14Micro Etchant14
7.1 7.2 8 9.1 9.2 9.3 9.4 9.5 9.6 9.7 9.8 9.9 9.1	Application Methods13Types.13TROUBLESHOOT GUIDE13GLOSSARY14Abrasion14Charging14Charging14Coupon Test Strip.14Crescent Moon Scratch14Grind.14Grinding Wheel14Grind Mount Holder14Micro Etchant14Mounting14
7.1 7.2 8 9.1 9.2 9.3 9.4 9.5 9.6 9.7 9.8 9.9 9.10 9.11	Application Methods13Types13TROUBLESHOOT GUIDE13GLOSSARY14Abrasion14Charging14Coupon Test Strip14Crescent Moon Scratch14Grind14Grind Wheel14Grit Size14Grind Mount Holder14Mounting14Polish Mount Holder14
7.1 7.2 8 9.1 9.2 9.3 9.4 9.5 9.6 9.7 9.8 9.9 9.1 9.1 9.1	Application Methods13Types13TROUBLESHOOT GUIDE13GLOSSARY14Abrasion14Charging14Coupon Test Strip14Crescent Moon Scratch14Grind14Grind Wheel14Grit Size14Grit Size14Micro Etchant14Mounting14Polish Mount Holder142Polish14
7.1 7.2 8 9.1 9.2 9.3 9.4 9.5 9.6 9.7 9.8 9.9 9.10 9.11	Application Methods13Types13TROUBLESHOOT GUIDE13GLOSSARY14Abrasion14Charging14Coupon Test Strip14Crescent Moon Scratch14Grind14Grinding Wheel14Grind Mount Holder14Micro Etchant14Polish Mount Holder14Sample Removal14
7.1 7.2 8 9.1 9.2 9.3 9.4 9.5 9.6 9.7 9.8 9.9 9.10 9.11 9.11 9.11	Application Methods13Types13TROUBLESHOOT GUIDE13GLOSSARY14Abrasion14Charging14Coupon Test Strip14Crescent Moon Scratch14Grind14Grind Wheel14Grit Size14Grind Mount Holder14Mounting14Polish Mount Holder14Abrasion14Crescent Moon Scratch14Grind Size14Grind Size14Grind Mount Holder14Mounting14Abrasion14Sample Removal14A Scratch14
7.1 7.2 8 9.1 9.2 9.3 9.4 9.5 9.6 9.7 9.8 9.9 9.10 9.11 9.12 9.13	Application Methods13Types13TROUBLESHOOT GUIDE13GLOSSARY14Abrasion14Charging14Coupon Test Strip14Crescent Moon Scratch14Grind14Grind Wheel14Grit Size14Grind Mount Holder14Micro Etchant14Polish Mount Holder14Sample Removal14Scratch14Scratch Trace14
7.1 7.2 8 9.1 9.2 9.3 9.4 9.5 9.6 9.7 9.8 9.9 9.1 9.1 9.1 9.1 9.1 2 9.1 2	Application Methods13Types.13TROUBLESHOOT GUIDE13GLOSSARY14Abrasion14Charging14Coupon Test Strip.14Crescent Moon Scratch14Grind14Grind Wheel14Grind Mount Holder14Micro Etchant14D Mounting14Polish Mount Holder14Sample Removal14Scratch Trace14Speed Bump14

Target Plated-Through Hole	15
Tooling Edge, Mount	15
Tooling Holes, Microsection	15
Tooling Pins, Microsection	15
REFERENCES	15
RECOMMENDED READING	15
ILLUSTRATIONS	15
	Tooling Edge, Mount Tooling Holes, Microsection Tooling Pins, Microsection REFERENCES RECOMMENDED READING

Figures

Figure 1a	Planar distortion	. 16
Figure 1b	Center line integrity	. 16
Figure 2	Target holes in same axis	. 17
Figure 3	Gaps between mounting material and samples	. 17
Figure 4	Depression in mounting material	. 17
Figure 5	Effects of mechanical grinding and polishing	. 18
Figure 6	Perpendicularity of motor to sandpaper disc assembly	. 18
Figure 7	Position of mount holder on sandpaper disc	. 19
Figure 8	Reference Zero relationship between target holes and tooling edge	. 19
Figure 9	Mount Holder Collar: Effects of Deflection	. 20
Figure 10	Effect of wear depression on mount holders	. 20
Figure 11	Balanced Mount Holder	. 21
Figure 12	Carbide pad flatness and height	. 21
Figure 13	Carbide pad should be center on arc	. 22
Figure 14	Leaf cut effect	. 22
Figure 15	Abrasive paper grit size (American vs. European)	. 23
Figure 16	Effect of trash trapped between mount holder and tooling edge	. 24
Figure 17	a Acceptable polish quality	. 25
Figure 17	b Acceptable polish quality	. 25
Figure 18	a Samples above potting material	. 25
Figure 18	b Samples above potting material	. 25
Figure 19	a Rounding at copper plate and solder interface	. 26
Figure 19	b Rounding at copper plate and solder interface on the surface	. 26
Figure 20	Scratch deformation	. 26
Figure 21	Smeared metal	. 26
Figure 22	Contact area comparison – sandpaper vs. polish cloth	. 27
Figure 23	-	

Tables

Table 1	Mounting Material Characteristics	. 5
Table 2	Etchant Types	14
	Troubleshoot Guide	28

Guidelines for High Volume Microsection

1 SCOPE

High volume microsection is a process. These guidelines discuss the many variables and problems associated with the process from sample removal to micro-etch. The guidelines do not promote any one vendor's process, but discuss the variables common to high volume microsection.

The process variables and problems are organized so the reader can research a specific issue or overview the variables of a process area.

2 APPLICABLE DOCUMENTS

2.1 IPC

IPC-D-275 Design Standard for Rigid Printed Boards and Rigid Printed Board Assemblies

IPC-T-50 Terms and Definitions for Interconnecting and Packaging Electronic Circuits

3 SAMPLE REMOVAL PROCESS

3.1 Sample Location

3.1.1 Coupon Test Strip Companies generally use a "home grown" or military conformance coupon for microsection inspection. IPC-D-275 outlines the attributes a coupon test strip should exhibit based on the product type being built.

Benefits:

- Production parts are not lost due to microsection testing.
- The internal and external features are the same from panel to panel to facilitate SPC data collection.
- The strips may be used to screen product as required.
- The customer can correlate to your microsection results easier because you both sample in the same location on the same test design.

Drawbacks:

- Space is lost on the panel that could be used to build parts.
- The test strip may not be representative of the associated part.

3.1.2 Part The actual production parts are used for microsection inspection.

Benefits:

• Space is not wasted on the panel due to test strips.

- There are no paneling constraints that dictate where the test strip must be placed to preserve part correlation.
- There is less of an issue over how representative the test strip is to the associated part.

Drawbacks:

- Microsection inspection of parts may not be cost effective for product with a high unit cost.
- For MLBs, multiple samples are usually microsectioned to inspect all the inner layer connections for each panel. These multiple samples can significantly increase the sample plan.
- The test results may not agree with the customer's results because microsections were taken on different locations of the part. This can only be resolved by providing the part sample locations to the customer.

3.2 Removal Method Regardless the method chosen, the cutting edge should remain a minimum of 0.25 cm [0.100 in] from the edge of the target plated-through-hole (PTH) pads. This is to prevent cutting deformation causing damage to the sample which may lead to false failures. The only exception to this guideline is abrasive cut-off wheels.

3.2.1 Fracturing This method is usually used in conjunction with routing to remove samples from brittle material (i.e., polyimide). The sample is routed leaving a finger tab that holds the sample in the panel. Fracturing is when the operator pushes or cuts the sample out of the panel by breaking the finger tab. The benefits and drawbacks of routing will be discussed in that section.

Benefits:

- The samples are routed and remain with the panel. This resolves panel traceability issues when the actual sample is not serialized.
- The samples, test strip, and parts are routed at one time. This prevents unnecessary use of costly production routers to only rout the sample.

Drawbacks:

- The finger tabs width needs to be optimized to keep the sample in the test strip during handling and permit an operator to push the sample out. The tab width may be different for families of products and/or board thickness. Thick boards may require needle nose pliers (or equivalent) to break the finger tabs.
- The location of the tab needs to be as far from the target PTHs and microsection tooling holes as possible. This will minimize the likelihood that material stresses will be transferred to the sample when it is pushed out.