



**Inks, colors, and supplies...your way.
Guaranteed!**

CERTIFICATION OF COMPLIANCE

1. Baby Soft Series Ink manufactured by One Stroke Inks
2. Section 108 of the Consumer Product Safety Improvement Act of 2008 including Subsection 108(a), Subsection 108(b)(1), Subsection 108(b)(2)
3. One Stroke Inks, 458 Roberts Ave, Louisville, Ky 40214, (502) 366-1070
4. Roger Bihl, 458 Roberts Ave, Louisville, Ky 40214, (800) 942-4447
5. Manufactured in January 2009 by One Stroke Inks, Louisville, Ky, USA
6. Tested on February 11, 2009 at IMR Test Labs
7. IMR Test Labs
131 Woodsedge Drive
Lansing Business Park
Lansing, NY 14882
(607) 533-7000
www.imrtest.com



131 Woodsedge Drive
Lansing Business & Technology Park
Lansing, NY 14882

Phone 607.533.7000 • Fax 607.533.9210
www.imrtest.com • E-mail imr@imrtest.com
Toll Free 888.464.8422

TEST REPORT

IMR Report Number 200901361P

March 18, 2009

Rocky Bihl
One Stroke Inks
458 Roberts Avenue
Louisville, KY 40214

PO Number
0032374

Date Received
February 11, 2009

Sample ID
Baby Soft Series Ink

Specification(s)
Consumer Product Safety
Improvement Act of 2008
(H.R. 4040, Title I, section
101, paragraph a)

SUMMARY

One ink sample was received for total lead analysis in accordance with the requirements of the Consumer Product Safety Improvement Act of 2008.

Based on the results, the sample **complies** with the lead requirement set forth in H.R. 4040, Title I, section 101, paragraph a.

The results are below.

RESULTS

Sample ID	Lead	Requirement ¹
Baby Soft Series Ink	<50	600 Maximum

Results in parts per million (ppm).

¹After 8/14/09, the requirement becomes 300 ppm maximum. After 8/14/11 the requirement becomes 100 ppm maximum.

Method(s): CAP-017J (ICP-AES)



Reviewed by

Brian Wackowicz
Lead Chemist

Reviewed by

Andrew J. Waldron
Lead Chemist

All procedures were performed in accordance with the IMR Quality Manual, current revision, and related procedures; and the PWA-MCL Manual F-23 and related procedures. The information contained in this test report represents only the material tested and may not be reproduced, except in full, without the written approval of IMR, Inc. IMR, Inc. maintains a quality system in compliance with the ISO/IEC 17025:2005 and is accredited by the American Association for Laboratory Accreditation (A2LA), certificates #1140.01 and #1140.02. IMR, Inc.'s liability to the customer or any third party is limited to the amount charged for services provided. All samples will be retained for a minimum of 6 months and may be destroyed thereafter unless otherwise specified by the customer. The recording of false, fictitious, or fraudulent statements or entries on this document may be punished as a felony under federal statutes.