



Michael R. Pence  
Governor

Jerome M. Adams, MD, MPH  
State Health Commissioner

**DATE:** December 2, 2015

**TO:** All Local Health Departments  
Attn: Chief Food Inspection Officer

**FROM:** *Laurie Kidwell*  
Laurie Kidwell, RRT Supervisor  
Food Protection Program

**SUBJECT:** Bestmed, LLC - RECALL [Medical Device]

**AFFECTED PRODUCT:** Digital Temple Thermometer

**SUMMARY:** Unclassified Recall; The recall was initiated after becoming aware of the inaccurate temperature readings by the Digital Temple Thermometer, and confirming the source of the performance problem with the manufacturer, K-Jump Health Co., Ltd.

The following model number(s) have been recalled:

KD-2201  
(Note: Model KD-2201L is not subject to this recall action)

The Digital Temple Thermometer is a hand-held thermometer and can be identified by the DTT™ logo on the face, as shown below:

The back of the thermometer has a label with the KD-2201 designation as shown below. The slash-mark (/) is not part of the model number. The lot number may be found on the bottom of the back label, immediately after the designation "S/N:". In the picture below, the lot number is S/N: 3113.

**SUGGESTED ACTION:** For consumer inquiry only. To arrange for the return and replacement of the thermometer or to ask questions, consumers should contact the Bestmed via telephone at (877) 299-6700, facsimile at (303) 271-0163, or email at [DTTRecall@bestmedusa.com](mailto:DTTRecall@bestmedusa.com), at any time.

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### Recall -- Firm Press Release

FDA posts press releases and other notices of recalls and market withdrawals from the firms involved as a service to consumers, the media, and other interested parties. FDA does not endorse either the product or the company.



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To promote and provide  
essential public health services.

*Bestmed, LLC Issues Nationwide Recall Of Digital Temple Thermometer (DTT™), Model No. KD-2201  
Manufactured By K-Jump Health Co., Ltd.*

FDA posts press releases and other notices of recalls and market withdrawals from the firms involved as a service to consumers, the media, and other interested parties. FDA does not endorse either the product or the company.

For Immediate Release

November 30, 2015

Contact

## Consumers/Media

Bestmed

[DTTRecall@bestmedusa.com](mailto:DTTRecall@bestmedusa.com)

(877) 299-6700

Fax: (303) 271-0163

Firm Press Release

[View Product Photos](#)

On November 12, 2015, Bestmed, LLC, a medical device distributor, initiated a nationwide recall of the Digital Temple Thermometer Model No. KD-2201 manufactured by K-Jump Health Co., Ltd, featuring lot numbers S/N: 3612 through S/N: 3715, which were sold between October 2012 until the start of the recall in November 2015.

Some Digital Temple Thermometers contain a manufacturing problem causing the affected thermometers to display temperatures that are inaccurate and lower than actual body temperatures, which potentially may cause the user or caregiver of the user to delay or forego seeking appropriate care (generally an over-the-counter fever reduction medication) or receive more care than appropriate, when relying solely on the temperature display on the thermometer.

Consumers who have a Digital Temple Thermometer should immediately stop using the device. Consumers may return the thermometer to Bestmed for a replacement thermometer.

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KD-2201

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Bestmed voluntarily initiated the recall after becoming aware of the inaccurate temperature readings by the Digital Temple Thermometer, and confirming the source of the performance problem with the manufacturer, K-Jump Health Co., Ltd.

The Digital Temple Thermometer was distributed throughout the United States for direct sale and resale through consumer retail stores, and through consumer retail stores in Canada under the following packaging labels:



1. Bestmed
2. Good Neighbor
3. Kroger
4. Medline
5. Meijer
6. Premier Value
7. Safeway
8. Life Brand
9. Target
10. Top Care
11. Best Choice (Value Merchandise)
12. Western Family

To arrange for the return and replacement of the thermometer or to ask questions, consumers should contact the Bestmed via telephone at (877) 299-6700, facsimile at (303) 271-0163, or email at [DTTRecall@bestmedusa.com](mailto:DTTRecall@bestmedusa.com), at any time. Our office is available Monday through Friday 9:00 AM to 7:30 PM, Eastern Time to speak with a consumer support specialist, and all messages and email will be responded to promptly.

Adverse reactions or quality problems experienced with the use of this product may be reported to the FDA's MedWatch Adverse Event Reporting program either online, by regular mail or by fax:

- Complete and submit the report **Online:** [www.fda.gov/medwatch/report.htm](http://www.fda.gov/medwatch/report.htm)
- **Regular Mail or Fax:** Download form [www.fda.gov/MedWatch/getforms.htm](http://www.fda.gov/MedWatch/getforms.htm) or call 1-800-332-1088 to request a reporting form, then complete and return to the address on the pre-addressed form, or submit by fax to 1-800-FDA-0178

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#### Product Photos



