June 9, 2003

ADDITIONAL LIPITOR LOTS RECALLED

On May 22, you were advised that certain lots of Lipitor 10 mg, NDC 0071-0155-23, which were repackaged by Med-Pro, Inc. of Lexington, NE were being voluntarily recalled.

In conjunction with the Food and Drug Administration, we are expanding our original voluntary recall to include all lots of Lipitor that Albers purchased, which were packaged by Med-Pro. Although the FDA has not Tested all of the additional lots, Albers is voluntarily recalling them as a precautionary measure. All lots subject to this recall are on the attached list.

The same procedures outlined to you on May 22, 2003 apply. You should immediately cease distribution or use of these products. This recall goes to the consumer level.

If you have distributed or sold any of these products to other distributors, retail pharmacies, or individual persons you should provide notice to them of this recall and advise them that the recall information should be provided to all consumers who received these products. These products should not be used. They should be returned immediately.

Products should be shipped to

Albers Medical Distributors, Inc

4400 Broadway, Suite 116

Kansas City, MO 64111

10 MG 90 TABLETS NDC 00071-0155-23

LOT NUMBER	EXPIRATION DATE
12912V	10/04
12912V 1	10/04
12912V 2	10/04
12912V 3	10/04
04132V 1	01/04
04132V 2	01/04
04132V 3	01/04
04132V 4	01/04
04132V 5	01/04
04132V 6	01/04
04132V 7	01/04
10772V 1	05/04
10772U 1	05/04
12542V 1	05/04
2072V 1	09/04
20722V 1	09/04
20722V 2	09/04
20722V 3	09/04
20722V 4	09/04
16942V	09/04
16942V 1	09/04
16942V 2	09/04
16942V 3	09/04
16942V 4	09/04
437023 1	02/05

20 MG 90 TABLETS NDC 00071-0156-23

LOT NUMBER	EXPIRATION DATE
027049D	02/04
D27048 1	02/04
D27049 1	02/04
511022 1	06/04
511022 2	06/04
511022 3	06/04