

**BILL SUMMARY**  
2<sup>nd</sup> Session of the 57<sup>th</sup> Legislature

<b>Bill No.:</b>	<b>HB 3418</b>
<b>Version:</b>	<b>Proposed Committee Substitute</b>
<b>Request Number:</b>	<b>10874</b>
<b>Author:</b>	<b>McEntire</b>
<b>Date:</b>	<b>2/25/2020</b>
<b>Impact:</b>	<b>Please see previous summary of this measure</b>

**Research Analysis**

The PCS to HB 3418 exempts the distribution of dialysate or peritoneal dialysis devices used for treating end-stage renal disease (ESRD) from the Oklahoma Pharmacy Act. However, dispensing facilities must ensure that the products are FDA-approved, held by a licensed manufacturer, delivered in their original packaging, delivered pursuant to a prescription, and delivered directly to the patient or health care provider responsible for administering the dialysis therapy. Products do not need to be certified by a pharmacist prior to delivery.

Prepared By: Anna Rouw

**Fiscal Analysis**

The measure is currently under review and impact information will be completed.

Prepared By: Mark Tygret

**Other Considerations**

None.