RESOLVED, That the American Bar Association urges states, territories and tribes to support the removal of legal barriers to the appropriate use by health care providers of Expedited Partner Therapy (EPT), applied as specified in protocols promulgated by the U.S. Centers for Disease Control and Prevention, in the treatment of those sexually transmitted diseases identified in the evidence-based recommendations of the CDC and the policy statements of the American Medical Association (adopted June 2006).
I. BACKGROUND AND INTRODUCTION

Despite major advances and achievements in the detection, treatment, and prevention of sexually transmitted diseases (STDs) in the United States, infections such as chlamydia and gonorrhea remain significant public health challenges. The U.S. Centers for Disease Control and Prevention (CDC) estimates that over 700,000 new cases of gonorrhea and 2.8 million new cases of chlamydia occur each year. Traditional practices to inform, evaluate and treat sex partners of persons infected with STDs have relied upon patients or health care providers to notify partners of infected persons of their exposure to an STD.

Initially developed to help control syphilis, partner management became widely recommended for gonorrhea, chlamydial infection and, most recently, human immunodeficiency virus (HIV) infection. However, for STDs other than syphilis, partner management based on provider referral is rarely assured, while patient referral has had only modest success in assuring partner treatment. An alternative approach to assuring treatment of partners is expedited partner therapy (EPT). EPT is the delivery of medications or prescriptions by persons infected with an STD to their sex partners without clinical assessment of the partners. Clinicians (e.g. physicians, nurse practitioners, physician assistants, pharmacists, public health workers) provide patients with sufficient medications directly or via prescription for the patients and their partners. This practice is intended to reduce the incidence of persistent or recurrent infection associated with cases where the STD-infected patient is engaging in ongoing sexual activity with partners whose concurrent infection is preventing adequate treatment, and is traditionally used in conjunction with physician guidance to patients to notify their sex partners of the infection.

The CDC produced an evidence review and associated guidance in 2006 on the use of EPT as an option for partner management for selected STDs and patients, based on 1) an internal review of scientific biomedical and behavioral evidence and 2) consultations including internal and external expertise from researchers, STD program managers, health departments, professional medical and public health organizations, and federal colleagues. CDC’s guidance indicates that EPT is a “useful option to facilitate partner management among heterosexual men and women with chlamydial infection or gonorrhea.” Additionally, the American Medical Association (AMA) adopted policy in June, 2006 that supported the CDC’s guidance on EPT as stated in the

1 The national Centers for Disease Control and Prevention (CDC) uses the terminology sexually transmitted diseases (STD). To be consistent with CDC language, the acronym STD will be used in this document.
5 Id. at 10.
6 Id. at 9.
7 Id. at 6.
CDC White Paper. The evidence-based recommendation of the CDC is that EPT is a useful option for the treatment of gonorrhea and chlamydia, but not for other STDs such as syphilis and trichomoniasis (see Section IV below).

EPT is typically a single-dose therapy with a broad-spectrum antibiotic. Where EPT is practiced, it can be done by giving the patient a dose for the patient’s partner, or giving the patient a prescription for enough pills for both the patient and partner. It is possible that the patient could also give the partner’s name (or a false name) for the prescription to be written by the physician for the partner.

Structured implementation of EPT according to CDC recommendations requires meeting various administrative requirements (e.g., prescription requirements, insurance and other payment issues), but the legality of EPT is largely unknown, even to members of groups traditionally attributed with responsibility for the legality of the practice. In order for EPT to reach its full potential as a treatment mechanism for chlamydial infection or gonorrhea in accordance with CDC guidelines, the legality of the practice must be clarified and any statutory impediments removed. To that end, the CDC and the Center for Law and the Public’s Health at Georgetown and Johns Hopkins Universities collaborated to assess the legal framework regarding EPT, the results of which are posted on the following link: http://www.cdc.gov/std/ept/legal/default.htm.

II. CURRENT APPLICATIONS OF PATIENT-DELIVERED PARTNER THERAPY

Partner notification is a cornerstone practice used in the treatment of patients diagnosed with STDs and includes guidance for the infected patient to refer their sex partners for diagnosis and treatment of STD infection, yet fewer than 20 percent of persons diagnosed with gonorrhea or chlamydia are offered assistance in notifying their sex partners. Healthcare practitioners face the reality that despite referrals encouraging patients to have their sex partners tested and treated, many partners fail to receive adequate treatment and reinfection is a common scenario. For example, a 2000 study showed that physicians across the country dispensed additional medications for partners of persons infected with gonorrhea or chlamydial infection on a widespread but essentially unregulated basis. Approximately one-half of the physicians surveyed practiced EPT one or more times, and an estimated one-seventh had done so on a
frequent basis. Additionally, one study found that more than 25 percent of medical providers in New York City have reported “frequent use” of EPT when diagnosing gonorrhea or chlamydia.\textsuperscript{14}

Proponents of EPT believe the inadequacies of current partner notification and referral guidance, combined with the increased incidence of persistent STD occurrence in the United States, warrant increased utilization of EPT and incorporation into clinical and public health policies.\textsuperscript{15} Emily J. Erbelding, M.D., M.P.H. and Jonathan M. Zenilman, M.D., of Johns Hopkins University, deem the findings “a major advance for the control and prevention of STDs”\textsuperscript{16} and conclude that “the use of expedited approaches designed to circumvent traditional evaluation by a clinician increases the chance of the exposed partner’s receiving proper therapy and, most important, reduces the original partner’s risk of infection.”\textsuperscript{17} EPT programs should include an educational component for medical providers authorized to dispense medications; the practice of EPT is designed to limit liability issues, as medical providers either dispense or prescribe standard medications, i.e. no medications known to be associated with extreme side effects.

The practice of EPT can be a useful tool in reducing the high rates of new cases of both gonorrhea and chlamydia that occur each year.\textsuperscript{18} EPT is an increasingly attractive option for clinicians faced with large numbers of chlamydial infections, decreasing staff to assist with partner notification, and the difficulties of getting male partners to providers for an infection that generally has no symptoms and which has no adverse health outcomes for males. Scientific evidence indicates that the recurrence of STD infection can be better reduced by giving the index patient medication to deliver to their partner(s) than by giving the index patient a standard referral service.\textsuperscript{19} The University of Washington in Seattle reported that “the provision of chlamydia or gonorrhea treatment directly to patients’ sexual partners, without requiring the partners to visit a physician, significantly improved infection control in patients.”\textsuperscript{20} Existing data also suggests that patients treated with EPT are more likely to notify their partners in accordance with physicians’ guidance.\textsuperscript{21}

Based on studies reporting the success of EPT and a study outlining risk factors associated with failure to notify potentially infected sexual partners, Public Health-Seattle and King County (PHSKC) in Washington began a partner notification program in 2004 to treat cases of gonorrhea and chlamydia.\textsuperscript{22} The program’s main features included the routine use of partner-delivered patient therapy by medical providers treating heterosexual individuals with gonorrhea or

\textsuperscript{15} Golden, \emph{supra} note 12, at 685.
\textsuperscript{16} Emily J. Erbelding, M.D., M.P.H. and Jonathan M. Zenilman, M.D., Toward Better Control of Sexually Transmitted Diseases, 352 NEW ENGLAND JOURNAL OF MEDICINE, 2005, 720-721, at 721.
\textsuperscript{17} See \emph{supra} note 16 at 720.
\textsuperscript{18} See \emph{supra} notes 2 and 3.
\textsuperscript{19} Pharmaceutical Journal Online News, Better Results for STIs if Patients Offered Treatment for Partners, 274 THE PHARMACEUTICAL JOURNAL, Feb. 19, 2005, at 199.
\textsuperscript{20} Kate Johnson, Patient-Delivered Treatment for Partners Reduces Chlamydia and Gonorrhea; Infectious Diseases; 7 FAMILY PRACTICE NEWS, Apr. 1, 2002, at 22
\textsuperscript{21} Hogben, \emph{supra} note 11 at 103.
chlamydia, and a case report forms designed to “triage patients at high risk for partner notification failure to receive public health partner notification assistance.” With respect to the effectiveness of the Washington program, researchers found:

2) The use of PDPT by providers in King County increased almost 3-fold concurrent with implementation of the program;
3) Patients who received PDPT from their provider were significantly less likely than other recently infected persons to report having untreated partners;
4) Providers successfully used the modified case report form to selectively refer patients with untreated partners to public health;
5) Patients referred to public health by providers usually accepted some form of partner management assistance; and
6) The estimated percentage of persons with gonorrhea or chlamydial infection for whom all partners were treated rose from 39% to 65% concurrent with the institution of the partner notification program.  

Thus, the findings of the randomized trial performed by PHSKC in Washington demonstrates the usefulness of EPT by medical providers as a tool to assist patients in partner notification and partner treatment, which has an inverse effect on gonorrhea and chlamydia morbidity rates.

Between 2000 and 2004, CDC sponsored four randomized controlled trials (RTC) designed to 1) compare EPT with standard partner management approaches and 2) assess behavioral predictors of treatment and reinfection. By late 2004, all trials had concluded data collection and analysis. The evidence across trials suggested EPT resulted in an approximate 20 percent reduction rate in recurrent or persistent infections among those originally diagnosed with chlamydial infection, and a 60 percent or better reduction rate among those originally diagnosed with gonorrhea. All trials reported favorable behavioral correlates, including increased notification and treatment of sex partners, and fewer instances of unprotected intercourse.

Following these findings, CDC convened two expert consultations in 2004 and 2005 to review scientific evidence related to EPT and address operational issues affecting its implementation. In February 2006, CDC issued a report providing the background of EPT, evidence in support of the practice, and guidance for using EPT as an option for managing partners of heterosexual sex partners with gonorrhea or chlamydial infection.24 In that report, CDC recommended EPT as a clinical option for heterosexual patients with gonorrhea or chlamydial infection. A trial based on EPT for women with trichomoniasis yielded null results and dubious behavioral correlates; CDC did not recommend routine use of EPT for patients diagnosed with trichomoniasis. CDC expects that partner management for syphilis be conducted with public health professional notification, where possible.

In 2008, the American Academy of Pediatrics passed the following resolution on EPT:

Resolved, that the Academy develop a policy statement that supports the practice of expedited partner therapy (EPT), i.e., treating the sex partners of sexually transmitted

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23 See supra note 22 at 602.
24 CDC Fact Sheet – Gonorrhea., supra note 2.
infections (STI)-infected persons without requiring the partners’ prior clinical evaluation as an alternative to traditional partner notification, and be it further Resolved, that the Academy support policies and support legislation that would allow a health care provider diagnosing *Chlamydia trachomatis* in an individual to prescribe or dispense antibiotics to that person’s sex partner(s) without examining them.

In June 2006, The American Medical Association (AMA) House of Delegates passed the following resolution in support of CDC’s guidance on the use of EPT:

The following statements, recommended by the Council on Science and Public Health, were adopted as by the AMA House of Delegates as AMA policy and directive at the 2006 AMA Annual Meeting:

1. The AMA supports the Centers for Disease Control and Prevention’s (CDC’s) guidance on expedited partner therapy (EPT) that was published in its 2006 white paper, *Expeditied Partner Therapy in the Management of Sexually Transmitted Diseases.* *(Policy)*

2. The AMA will continue to work with the CDC as it implements EPT, such as through the development of tools for local health departments and health care professionals to facilitate the appropriate use of this therapy *(Directive)*

In September 2006, CDC’s *STD Treatment Guidelines* endorsed EPT for the first time:

When medical evaluation, counseling, and treatment of partners cannot be done because of the particular circumstances of a patient or partner or because of resource limitations, other partner management options can be considered. One option is patient-delivered therapy, a form of expedited partner therapy (EPT) in which partners of infected patients are treated without previous medical evaluation or prevention counseling… Medications and prescriptions for patient-delivered therapy should be accompanied by treatment instructions, appropriate warnings about taking medications if pregnant, general health counseling, and advice that partners should seek personal medical evaluations, particularly women with symptoms of STDs or [pelvic inflammatory disease].25

EPT holds great potential as a tool in reducing the rate of persistent or recurrent STDs, and is touted as a major advance in the treatment of STDs. While EPT is not yet a widespread practice used by physicians in the treatment of STDs, it is used by some practitioners in both the private and public sectors. EPT is part of a host of guidance and advocacy tools, but it is still widely underused despite proven benefits in practice.

### III. BARRIERS TO GREATER IMPLEMENTATION OF EPT

There are a number of barriers to the widespread use of EPT in the treatment of STDs, some of which are medical in nature. For example, some physicians are more likely to utilize EPT when treating heterosexual female index patients rather than heterosexual male index patients because

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of a general reluctance to treat the former for STDs without first performing a pregnancy test or screening for pelvic inflammatory disease.\textsuperscript{26} In addition, EPT is contraindicated for STDs where the preferred treatment option is something other than an orally administered prescription, such as the treatment for syphilis.\textsuperscript{27} Other barriers include physicians’ concerns that the medication will not be delivered to their patients’ sex partners, accidental dispensing of multiple dosage, and missed opportunities to counsel partners.\textsuperscript{28}

However, one of EPT’s greatest barriers is legal in nature. The results of one study indicated that despite its potential benefits, an overwhelming 75 percent of physicians did not use EPT because of concerns about liability.\textsuperscript{29} According to a second study, 36 percent of primary care physicians do not use EPT out of fear of potential liability, even though approximately 90 percent agree that EPT helps them provide better care for their patients with chlamydia and protects their patients from reinfection.\textsuperscript{30} Physicians are also joined by nurse practitioners in their uncertainty: 28 percent fear using EPT because it may expose them to the possibility of a lawsuit, although 50 percent of nurse practitioners agree that the practice protects their patients from reinfection.\textsuperscript{31}

Physicians’ and nurse practitioners’ concerns are not without reason because the legal status of EPT is dishearteningly uncertain. For example, EPT is expressly forbidden in some states, while legal in others.\textsuperscript{32} In addition, medical practice statutes may explicitly prohibit physicians from prescribing medication in the absence of a physician-patient relationship, and pharmacy boards may impose similar restrictions.\textsuperscript{33} An additional study, primarily concerned with medical and pharmacy boards, demonstrated confusion regarding the legal status of EPT. In this study, three states’ medical and pharmacy board respondents disagreed with one another when polled about the legality of EPT in their states.\textsuperscript{34}

While further research may be performed to address many of the medical concerns facing EPT, the aforementioned legal concerns present a large measure of the barriers to greater use of EPT as a clinical tool in treating STDs that research alone cannot dispel. In order for EPT to achieve its potential as a treatment option, its legal barriers must be addressed.

IV. CDC GUIDANCE FOR THE APPROPRIATE IMPLEMENTATION OF EPT

As analyses of the data from the EPT RTCs were completed, CDC turned its attention to interpretation of the results and translating the research into clinical guidance. In 2005, CDC undertook a review of evidence and a broad-scope consideration of the practice’s limitations and

\textsuperscript{26} Hogben, supra note 11 at 103.
\textsuperscript{27} Erbelding and Zenilman, supra note 16 at 721.
\textsuperscript{28} Linda M. Niccolai, Ph.D. and Diana M. Winston, M.P.H., Physicians’ Opinions on Partner Management for Nonviral Sexually Transmitted Infections, 28 American Journal of Preventive Medicine, 2005, 229-233, at 229.
\textsuperscript{29} See supra note 28 at 230.
\textsuperscript{31} Id. at 461 (see Table 5).
\textsuperscript{32} Erbelding and Zenilman, supra note 19 at 721.
\textsuperscript{33} Id.
\textsuperscript{34} Golden, supra note 9 at 113.
benefits. The result of the review and consultation was a published white paper entitled “Expedited Partner Therapy in the Management of Sexually Transmitted Diseases: Review and Guidance.” The CDC white paper represented a thorough, systematic review of available (published and unpublished) literature reviewing EPT and patient-delivered partner therapy. The paper provided a detailed summary of concerns regarding the implementation of EPT on a larger scale, as well as precise guidance for the use of EPT in practice.

The medical issues highlighted in the CDC white paper covered the gamut of concerns that physicians and nurse practitioners express, as noted above. In addition, the white paper addressed a range of implementation barriers to EPT: concerns about the coverage of direct and indirect costs related to EPT and the limitations on third-party coverage; administrative and practical considerations; missed opportunities for counseling; the attitudes and beliefs of health care providers and agencies, and confidentiality concerns.

Significantly, the CDC has found that confidentiality concerns arising from a patient being called upon to communicate personal medical issues with sexual partners are not germane to EPT, but are relevant to all patient referral forms of partner management, which actually comprise the vast majority of partner treatment interactions in the U.S. In addition, the white paper highlighted the current legal uncertainty of EPT, as well as isolated statutory impediments that exist in certain states. Finally, the paper acknowledged a new barrier: that “[t]he medicolegal ramifications may be uncertain in the event of adverse outcomes in the recipients of EPT.”

The white paper summarized these barriers into a listing of “implementation issues” affecting the utilization of EPT and its priority with respect to other, more traditional partner management strategies. The implementation issues are:

- limited study focus on special populations,
- possible presence of other STDs,
- STD co-morbidity in sex partners,
- potential effects of drug use,
- adverse effects of drug use,
- missed opportunities for prevention counseling,
- the uncertain legal status of EPT,
- medicolegal concerns of the risk (or perceived risk) of increased litigation,
- funding,
- privacy,
- drug delivery and packaging,
- the providers’ and health agencies’ attitudes and beliefs,
- administrative barriers,

36 CDC White Paper, supra note 4 at 4.
37 CDC White Paper, supra note 4 at 6.
38 Id. at 6.
40 See supra note 4 at 6.
provider education,
interaction with other partner management strategies, and
implications on retesting for chlamydial infection and gonorrhea.\textsuperscript{41}

The white paper concludes that EPT is at least equivalent, if not better, than standard patient referral in preventing persistent or recurrent chlamydial or gonorrheal infection.\textsuperscript{42} With all of the aforementioned implementation issues considered, the CDC white paper provides clear guidance for the use of EPT in practice. The CDC’s recommendations are five-fold:

- **Gonorrhea and Chlamydial Infection in Women:** EPT can be used to treat the sex partners of female patients infected with gonorrhea or chlamydia when other management strategies are impractical or unsuccessful.

- **Gonorrhea and Chlamydial Infection in Men:** EPT can be used to treat the sex partners of male patients infected with gonorrhea or chlamydia when other management strategies are impractical or unsuccessful, provided that female recipients of EPT are strongly encouraged to seek medical attention (this is particularly important for female recipients exhibiting symptoms of acute pelvic inflammatory disease such as abdominal or pelvic pain) in addition to accepting therapy.

- **Gonorrhea and Chlamydial Infection in Men Having Sex with Men:** Because of a lack of statistically significant data regarding the efficacy of EPT in this population (because of high risk of co-morbid, undiagnosed HIV), EPT is not suggested as a routine therapy for the male sex partners of male patients infected with gonorrhea or chlamydia. EPT is only suggested for selective use, with caution, when other strategies prove impractical or unsuccessful.

- **Women with Trichomoniasis:** Because of a high-risk of STD co-morbidity in sex partners (especially gonorrhea and chlamydia infection), EPT is not advised for use as a routine therapy for female patients with trichomoniasis. EPT is only suggested for use selectively, with caution, when other strategies prove impractical or unsuccessful.

- **Syphilis:** EPT is not suggested for use in the treatment of patients with infectious syphilis. Syphilis typically requires injection therapy, and partner notification services are ordinarily available at local or state health departments.\textsuperscript{43}

In summary, the CDC concludes that EPT is a useful option in the facilitation of partner management, particularly for the treatment of male partners of women with gonorrhea or chlamydia and the treatment of female partners of men with gonorrhea or chlamydia, and recommends the use of EPT as a tool in the prevention of persistent or recurrent gonorrhea or chlamydia infection in the treatment of women diagnosed with gonorrhea or chlamydia, or in the treatment of heterosexual men diagnosed with gonorrhea or chlamydia.\textsuperscript{44} EPT is not recommended for use in treating chlamydia or gonorrhea in men who have sex with men because of high rates of undiagnosed HIV and other STDs. EPT is also not recommended for treatment of chlamydia or gonorrhea among homosexual women because no data currently exist.

\textsuperscript{41} Id. at 18-24.
\textsuperscript{42} Id. at 34.
\textsuperscript{43} CDC White Paper, supra note 4 at 6.
demonstrating the efficacy or role of EPT among this population. Finally, existing data support the practice of EPT in treating chlamydia and gonorrhea only, because other STDs frequently require more invasive treatment procedures or medications with higher rates of allergic reactions. For example, EPT would be inappropriate for use in treating syphilis because the disease frequently requires injection therapy and many patients have allergic reactions to penicillin.

These guidelines are supported by the American Medical Association, whose House of Delegates passed a supporting resolution in June, 2006. The AMA’s recommendation adopts the CDC guidelines and pledges to support the CDC in the implementation of appropriate use of EPT. The American Academy of Pediatrics adopted a resolution in 2008 encouraging the practice of EPT and supporting policies and legislation allowing physicians to dispense medication for chlamydia to a person’s sexual partners without first examining them. The next step in addressing the implementation issues outlined in the CDC white paper is to address legal barriers to the use of EPT.

V. STATE-LEVEL LEGAL ANALYSIS OF THE LEGAL STATUS OF EPT

CDC and the Center for Law and the Public’s Health at Georgetown and Johns Hopkins Universities (Center) collaborated to assess the state-level legal framework concerning EPT to assist state and local STD programs in their efforts to implement EPT as an additional partner management tool. The primary research objective was to identify legal provisions that impact a clinician’s ability to provide a prescription for a patient’s sex partner, without prior evaluation of that partner, for purposes of treating an STD (specifically chlamydia or gonorrhea). Three broad legal frameworks were examined: 1) medical licensing and liability, 2) public health and safety, and 3) pharmaceutical practices. The examination of pharmaceutical practices focused on laws concerning drugs generally, and “legend” drugs (which require a prescription, such as most antibiotics), but not laws concerning controlled substances (e.g., habitual drugs regulated under the federal Controlled Substances Act). Antibiotics used to treat chlamydia and gonorrhea are not controlled substances.

In each of these three major areas, research included interpretive analysis of relevant laws and policies (i.e. statutes, bills, administrative regulations, judicial cases, administrative opinions) found through legal research engines (e.g., Westlaw, LEXIS), and publicly-available legal Websites. Secondary resources (e.g., reports, articles, media accounts) and informal discussions with federal, state, and local law and policy-makers, public health officials, and academics were used sparingly to gather some data. Information from these sources was confirmed through original legal research. Data were organized in a table that stratified references to relevant statutes, administrative regulations, cases, legislative bills, administrative orders, and medical or pharmaceutical board opinions. Tables are available in hard copy and on the CDC website at:

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45 CDC White Paper, supra note 4 at 33.
46 See supra note 4 at 33.
http://www.cdc.gov/std/ept/legal/default.htm. Included in the tables are hotlinks to relevant legal
documents analyzed and used to draw conclusions about the legal status of EPT in each state.

For each jurisdiction, one of three possible conclusions is offered, based on this assessment of
the multifarious laws and policies: 1) EPT is permissible (because the laws within the
jurisdiction either expressly allow the practice of EPT, or do not expressly prohibit it); 2) EPT is
possible (because the laws within the jurisdiction may allow EPT subject to specific
interpretations of inconsistent or amorphous provisions, supporting policies underlying legal
authorization, or incorporation by reference of current STD treatment guidelines); or 3) EPT is
likely prohibited (because the laws within the jurisdiction do not support the practice of EPT by
clinicians or others).49 The findings indicate that EPT is permissible or possible in a majority of
jurisdictions. At present, EPT is permissible in 10 jurisdictions, possible in 29 jurisdictions, and
likely prohibited in 13 jurisdictions.50

One question that arises is whether EPT might present confidentiality or privacy concerns for the
partner, especially since the underlying premise of EPT is that the partner may not know of the
infection or has chosen not to visit the doctor to be treated. EPT need not be invasive of the
privacy of the partner. For example, it is possible to practice EPT by providing the antibiotic
drug directly to the patient for use by the partner, or a prescription to the patient for a dose that is
adequate for both patient and partner. The antibiotic is broad-spectrum antibiotic and thus in
itself should not label the patient or partner as having or potentially having an STD, even if the
prescription were written to the partner’s name. Since STDs are required to be reported by
physicians to the state health department under state laws, EPT should not be any more invasive
of privacy than current public health practice.

As noted above, the threat of medical malpractice liability might pose a barrier to EPT. These
legal issues would need to be addressed by states, such as through adoption of protocols on how
to appropriately implement EPT.

VI. CASE STUDY: LEGAL IMPLEMENTATION OF EPT IN MARYLAND

Senate Bill 349 was signed into law on April 24, 2007, establishing a three-year Expedited
Partner Therapy Pilot Program (Program) in the Baltimore City Health Department. The purpose
of the Program is “to provide antibiotic therapy to a partner of a patient diagnosed with a
sexually transmitted infection of gonorrhea or chlamydia in order to contain the infection and
prevent further transmission.”51 Under the Program, medical providers, including licensed
physicians, certified nurse practitioners, and physician assistants, may dispense antibiotic therapy
for the partners of patients presenting with gonorrhea or chlamydia without a prior physical
examination.52 The Fiscal and Policy Note accompanying the Bill specifically cited CDC’s
inclusion of EPT in the 2006 Sexually Transmitted Diseases Treatment Guidelines as providing
validity for the practice of EPT.53 In addition, the Fiscal and Policy Note discussed the “Legal

49 Hodge, supra note 47 at 5.
50 See supra note 47 at 5.
51 MD. CODE REGS. 10.06.01.17-1 (2007)
52 Id.
53 http://mlis.state.md.us/2007RS/fnotes/bil_0009/sb0349.pdf
Assessment Concerning Expedited Partner Therapies” produced jointly by NCHHSTP/CDC and the Center for Law and the Public’s Health as legal support for EPT’s practice.\textsuperscript{54} The Program, which took effect on July 1, 2007, will remain effective for 3 years.\textsuperscript{55} At the end of each calendar year, the Baltimore City Health Department must submit a report to the Governor and the General Assembly detailing the “operation and performance” of the Program.\textsuperscript{56, 57}

\section*{VII. ABA COLLABORATION WITH CDC}

The ABA works diligently in addressing innovative issues in health law. In addition, the ABA has an ongoing collaboration with the CDC as documented in a Memorandum of Understanding signed by the two organizations.\textsuperscript{58} The Memorandum of Understanding joins the ABA and CDC in a mission to address issues relating to public health and the law. This resolution is a step that furthers that mission.

\begin{thebibliography}{9}
\bibitem{54} Id.
\bibitem{55} \textsc{Md. Code Regs. 10.06.01.17-1} (2007)
\bibitem{56} \textsc{Md. Code Ann., [Health-General] § 18-214.1(f)} (West 2007)
\bibitem{57} The report for this program is not publically available.
\bibitem{58} Memorandum of Understanding between the Centers for Disease Control and Prevention and the American Bar Association, signed March 7, 2005, \textit{available at} http://www2a.cdc.gov/phlp/docs/ABA.CDC.MOU.pdf (last visited January 11, 2007).
\end{thebibliography}
VIII. CONCLUSIONS

EPT is a useful tool in the fight against the increasing incidence of persistent and recurrent gonorrhea and chlamydia infection in the United States. Evidence substantiates the safety and usefulness of EPT in practice, and the CDC white paper addresses continuing issues to implementation. These issues encompass a number of medical and legal concerns faced by physicians, nurse practitioners, and pharmacists in the implementation of EPT. The CDC’s white paper proposes a clear set of guidelines for the use of EPT in practice today. This guidance allows for the use of EPT in the treatment of female sex partners of men and the male sex partners of women infected with gonorrhea or chlamydia. The AMA’s adoption of the CDC’s guidance and its and the American Academy of Pediatrics’ support in the implementation of EPT will address many of the medical issues hindering EPT’s widespread use by physicians and nurse practitioners, but cannot address the legal barriers.

Those in the legal profession have an opportunity to address the legal issues hindering EPT’s potential to reduce the rates of recurrent and persistent infections of gonorrhea and chlamydia. The ABA’s unique collaborative relationship with CDC makes the joint Memorandum of Understanding the prime vehicle for exploring the legal barriers to implementation of EPT on a larger scale, the success for which is dependent upon ABA support.

Therefore, the recommendations set forth above are consistent with the ABA’s missions and goals, in particular Goals III and IV, respectively, “To provide ongoing leadership in improving the law to serve the changing needs of society” and “To increase public understanding of and respect for the law, the legal process, and the role of the legal profession.”

Respectfully submitted,

Andrew J. Demetriou
Chair
Health Law Section
American Bar Association

August 2008

GENERAL INFORMATION FORM

To Be Appended to Reports with Recommendations
(Please refer to instructions for completing this form.)

Submitting Entity: Health Law Section
Submitted By: Andrew J. Demetriou, Chair

1. Summary of Recommendation(s).

The report and recommendation urges the American Bar Association to support the elimination of legal barriers to the provision by healthcare providers of Expedited Partner Therapy (EPT) in the treatment of certain sexually transmitted diseases identified by the U.S. Centers for Disease Control and Prevention and the policy statements of the American Medical Association on this issue, and to address any potential medico-legal implications posed by the practice of EPT.

2. Approval by Submitting Entity.

Approved by Section Council May 6, 2008.

3. Has this or a similar recommendation been submitted to the House or Board previously?

No.

4. What existing Association policies are relevant to this recommendation and how would they be affected by its adoption?

There are no Association policies addressed by this recommendation.

5. What urgency exists which requires action at this meeting of the House?

The need to address the over 700,000 new cases of gonorrhea and 2.8 million cases of chlamydia that occur each year makes action needed as soon as possible.

6. Status of Legislation. (If applicable.)

None pending.

7. Cost to the Association. (Both direct and indirect costs.)

None.
8. Disclosure of Interest. (If applicable.)

None.

9. Referrals.

This report and recommendation has been referred to the Sections of Individual Rights and Responsibility, Science & Technology, Litigation, Torts, Trial and Insurance Practice, and Administrative Law, and the Young Lawyers and General Practice, Solo and Small Firm Divisions, and the Standing Committee on Medical Professional Liability.

10. Contact Person. (Prior to the meeting.)

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11. Contact Person. (Who will present the report to the House.)

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Executive Summary

Summary of Recommendation

The report and recommendation urges the American Bar Association to support the elimination of legal barriers to the provision by healthcare providers of Expedited Partner Therapy (EPT) in the treatment of certain sexually transmitted diseases identified by the U.S. Centers for Disease Control and Prevention and the policy statements of the American Medical Association on this issue, and to address any potential medico-legal implications posed by the practice of EPT.

Summary of issues which the recommendation addresses

The recommendation calls for States, Territorial and Tribal governments to eliminate laws that would prohibit doctors from utilizing this treatment option, consistent with recommendations of the US CDC.

Explanation of how the proposed policy position will address the issue

The policy will encourage states, tribes and territories to eliminate statutory barriers to implementing the Expedited Partner Therapy (EPT)

Summary of minority views or opposition which have been identified

No opposing views have been identified.